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(54) Title: ANTIGENIC POLYPEPTIDES

(57) Abstract: The invention relates to a method for the identification of antigenic polypeptides, typically opsonic antigens, ex-
pressed by pathogenic microbes; vaccines comprising said antigens; and therapeutic antibodies directed to said antigenic polypep-
tides.

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INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07K7/04 C07K14/195 C07K16/12 A61K39/02 A61P31/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, BIOSIS, EMBASE, EMBL, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE EMBL [Online] 16 March 1999 (1999-03-16), BARASH ET AL: "Staphylococcus aureus polynucleotides and sequences" XP002250642 retrieved from AAW89789 accession no. EBI Database accession no. AAW89789 * Refers to EP-A-786519, published 30.07.97 (3271 pages); identical with Locus 1, Sequence 3 [4-363 : 2-361]; and SEQ 544 (EP), complete reversed DNA overlap [1400-5088 : 3689-1/Locus 1] *</p> <p style="text-align: center;">----- -/--</p>	<p>1-7, 9-16, 18-26</p>

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the International filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

Inter al Application No
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE EMBL [Online] 1 June 2001 (2001-06-01), KURODA ET AL: "Whole genome sequencing of meticillin-resistant Staphylococcus aureus" XP002250643 retrieved from Q99W10 accession no. EBI Database accession no. Q99W10 * 98% overlap in the region 21-251 [Locus 1, Sequence 4] : 1-231; misfits at 49, 83,141,144 and 229 (of Q99W10) *</p>	1
P,X	<p>WO 01 98499 A (UNIVERSITY OF SHEFFIELD / BIOSYNEXUS) 27 December 2001 (2001-12-27)</p>	1-7, 9-16, 18-26
P,Y	<p>* See the whole document - antigenic polypeptides from Staphylococcus aureus; SEQ.ID. 32 = identical with Locus 1, Sequence 1; page 5 -> SEREX *</p>	27
Y	<p>SAHIN ET AL: "Serological identification of human tumor antigens" CURRENT OPINION IN IMMUNOLOGY, vol. 9, no. 5, October 1997 (1997-10), pages 709-716, XP004313590 ISSN: 0952-7915 * The original SEREX method / see page 5 of the Application *</p>	27
A	<p>US 6 159 469 A (CHOI ET AL) 12 December 2000 (2000-12-12) * See Abstract - antigenic polypeptides from Streptococcus pneumoniae *</p>	1-26
A	<p>US 6 086 896 A (SPARLING ET AL) 11 July 2000 (2000-07-11) * See Abstract - antigenic polypeptide from Neisseria meningitidis *</p>	1-26
A	<p>US 5 543 323 A (RIDLEY ET AL) 6 August 1996 (1996-08-06) * See Abstract - antigenic polypeptides from Plasmodium *</p>	1-26
A	<p>WOOD ET AL: "Identification of antigenic sites on staphylococcal enterotoxin B and toxoid" FEMS IMMUNOLOGY AND MEDICINAL MICROBIOLOGY, vol. 17, 1997, pages 1-10, XP002250576 * See pages 8-9 (3.3 and 4) *</p>	1-26
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INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 02/03606

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
L	<p>DATABASE EMBL [Online] 20 February 2003 (2003-02-20), MASIGNANI ET AL: "Staphylococcus aureus proteins and nucleic acids" XP002250644 retrieved from AX618827 accession no. EBI Database accession no. AX618827 * Refers to W002094868, published 28.11.02 (international filing date 27.03.02, priority date 27.03.01) without sequences (electronically filed only) - see Locus 1, Sequence 1 = 100% identity *</p>	1-26
L	<p>DATABASE EMBL [Online] 20 February 2003 (2003-02-20), MASIGNANI: "Staphylococcus aureus proteins and nucleic acids" XP002250645 retrieved from AX618829 accession no. EBI Database accession no. AX618829 * As above; identical with Locus 1, Sequence 2 (except the first amino acid) *</p>	1-26
L	<p>DATABASE EMBL [Online] 20 February 2003 (2003-02-20), MASIGNANI: "Staphylococcus aureus proteins and nucleic acids" XP002250646 retrieved from AX618833 accession no. EBI Database accession no. AX618833 * As above; identical with Locus 1, Sequence 3 (except the first amino acid) *</p>	1-26
L	<p>DATABASE EMBL [Online] 20 February 2003 (2003-02-20), MASIGNANI: "Staphylococcus aureus proteins and nucleic acids" XP002250647 retrieved from AX618835 accession no. EBI Database accession no. AX618835 * As above; identical with Locus 1, Sequence 4 (except the first amino acid; erroneous omission of 241-251 ?) *</p>	1-26

INTERNATIONAL SEARCH REPORT

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Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; It is covered by claims Nos.:
1-26 (all partially) and 27 (entirely)

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although Claims 12-17 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the polypeptides/compositions.

Note also that "or part thereof" (Claim 1) has no clear meaning - it would even cover dipeptides in an extreme interpretation.

The applicant's attention is drawn to the fact that claims relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure. If the application proceeds into the regional phase before the EPO, the applicant is reminded that a search may be carried out during examination before the EPO (see EPO Guideline C-VI, 8.5), should the problems which led to the Article 17(2) declaration be overcome.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1-26 (all partially) and 27 (entirely)

Invention 1:

Claim 27 (the method used) and a first group of antigenic polypeptides (the 4 peptides of Locus 1, encoded by the first DNA sequence in Table 7), including their uses etc. as of dependent Claims 2-26, as applicable.

Inventions 2-134:

As invention 1 but limited to each subsequent group of peptides as encoded by the 2nd, 3rd, ..., 122th DNA sequence in Table 7, and the 123th, ..., 134th DNA sequence in Table 9, as applicable.

Note:

As a consequence of the lack of information in the Description about sequence relations (e.g. common subsequences ?) etc, the actual number of inventions may deviate from the above.

This is, however, not of significance at present.

INTERNATIONAL SEARCH REPORT

information on patent family members

Inte Application No
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Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0198499	A	27-12-2001	AU 7424801 A	02-01-2002
			BR 0111823 A	10-06-2003
			CA 2412504 A1	27-12-2001
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US 6159469	A	12-12-2000	US 6573082 B1	03-06-2003
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			AU 633306 B2	28-01-1993
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			DE 69004721 T2	17-03-1994
			DK 388738 T3	17-01-1994
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			JP 3047088 A	28-02-1991
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			ZA 9001757 A	28-11-1990

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AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
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KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
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ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

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ern Bank, Sheffield S10 2TN (GB). BRUMMEL, Kirsty

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For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
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(54) Title: ANTIGENIC POLYPEPTIDES

(57) Abstract: The invention relates to a method for the identification of antigenic polypeptides, typically opsonic antigens, ex-
pressed by pathogenic microbes; vaccines comprising said antigens; and therapeutic antibodies directed to said antigenic polypep-
tides.



WO 03/011899 A2

Antigenic Polypeptides

The invention relates to a method for the identification of antigenic polypeptides, typically opsonic antigens, expressed by pathogenic microbes; vaccines comprising
5 said antigens; and therapeutic antibodies directed to said antigenic polypeptides.

Microbial organisms cause a number of fatal or debilitating diseases which affect many millions of people around the world. Currently methods to control microbial organisms include the use of antimicrobial agents (antibiotics) and disinfectants.
10 These have proved to be problematic since exposure to these agents places a significant selection pressure resulting in the creation of resistant microbes which can avoid the effects of the antimicrobial agent(s). For example, recently it has been discovered that microbial organisms have become resistant to triclosan, an agent added to many disinfectants used in households and industrial environments.

15 An arguably greater problem is the evolution of antibiotic resistant strains of a number of significant pathogenic microbes.

For example, and not by way of limitation, it is estimated that there are up to
20 50 million people world-wide infected with drug resistant tuberculosis (TB) (Figures from the World Health Organisation, 1998). In the past the use of antibiotics to treat TB relied on the administration of single drugs (eg ethionamide) which promoted a relatively high frequency of resistance. For this reason, combinations of drugs are now used to treat tuberculosis. However the fatality rate in cases caused by strains
25 that are resistant to at least one drug used to treat tuberculosis still approaches 50% even when treatment is given. *Mycobacterium tuberculosis*, the causative agent of TB, is a slow growing bacteria and takes a long time to kill. Therefore, for a drug combination to be effective a person with TB must take the drug combination daily for at least six months. Accordingly, patients frequently have to take two or more
30 pills daily and this requires a regimented dosage over a relatively long period of treatment. Many patients take the medications only intermittently and therefore do

not finish the full course of therapy to completely eradicate the *M. tuberculosis* infection. Moreover, TB is strongly associated with HIV infection and therefore the establishment of TB is strongly correlated with immunosuppression.

- 5 Vaccination against TB has been available for many years. The bacillus calmette and guerin (BCG) vaccination has been widely used throughout the world for a long time because it is a safe and inexpensive means to vaccinate large numbers of people who potentially could contract TB. BCG is derived from live, attenuated strains of *Mycobacterium bovis*. However the impact of vaccination on the infectious forms of
- 10 TB is minimal and BCG has therefore contributed little to the overall control of the disease.

- A further example of a pathogenic organism which has developed resistance to antibiotics is *Staphylococcus aureus*. *S.aureus* is a bacterium whose normal habitat
- 15 is the epithelial lining of the nose in about 20-40% of normal healthy people and is also commonly found on people's skin usually without causing harm. However, in certain circumstances, particularly when skin is damaged, this germ can cause infection. This is a particular problem in hospitals where patients may have surgical procedures and/or be taking immunosuppressive drugs. These patients are much
- 20 more vulnerable to infection with *S.aureus* because of the treatment they have received. Resistant strains of *S.aureus* have arisen in recent years. Methicillin resistant strains are prevalent and many of these resistant strains are also resistant to several other antibiotics. Currently there is no effective vaccination procedure for *S. aureus*. In the US, *S.aureus* infections are the cause of 13% of the two million
- 25 hospitalised infections each year. This represents 260,000 people with an infection of *S.aureus*, of which 60-80,000 die.

- S. aureus* is therefore a major human pathogen capable of causing a wide range of life threatening diseases including septicaemia, endocarditis, arthritis and toxic
- 30 shock. This ability is determined by the versatility of the organism and its arsenal of components involved in virulence. Pathogenicity is multifactorial and no one

component has shown to be responsible for a particular infection, see Projan, S.J. & Novick, R.P. (1997) in *The Staphylococci in Human Disease* (Crossley, K.B. & Archer, G.L., eds.) pp.55-81.

5 At the onset of infection, and as it progresses, the needs and environment of the organism changes and this is mirrored by a corresponding alteration in the virulence determinants which *S. aureus* produces. At the beginning of infection it is important for the pathogen to adhere to host tissues and so a large repertoire of cell surface associated attachment proteins are made. These include collagen-, fibrinogen- and
10 fibronectin-binding proteins. The pathogen also has the ability to evade host defences by the production of factors that reduce phagocytosis or interfere with the ability of the cells to be recognised by circulating antibodies.

Often a focus of infection develops as an abscess and the number of organisms
15 increases. *S. aureus* has the ability to monitor its own cell density by the production of a quorum sensing peptide. Accumulation of the peptide, associated with physiological changes brought about by the beginning of starvation of the cells, elicits a switch in virulence determinant production from adhesins to components involved in invasion and tissue penetration. These include a wide range of
20 hemolysins, proteases and other degradative enzymes.

During the process of any infection the virulence determinants made by *S. aureus* are produced in response to environmental and physiological stimuli. These stimuli will be dependent on the niche within the body and will change as the infection
25 progresses. Little is known of the conditions *in vivo* and it is likely that some components are produced solely in this environment. These are therefore potential vaccine components, which could not be discovered by previous techniques.

One of the most important developments in recent medical history is the development of vaccines which provide prophylactic protection from a wide variety of pathogenic organisms. Many vaccines are produced by inactivated or attenuated pathogens which are injected into an individual. The immunised individual responds
5 by producing both a humoral (antibody) and cellular (cytolytic T cells, CTL's) response. For example, hepatitis vaccines are made by heat inactivating the virus and treating it with a cross linking agent such as formaldehyde. An example of an attenuated pathogen useful as a vaccine is represented by polio vaccines which are produced by attenuating a live pathogen.

10

However the use of attenuated organisms in vaccines for certain diseases is problematic due to the lack of knowledge regarding the pathology of the condition and the nature of the attenuation. For certain viral agents this is a particular problem since viruses, in particular retroviruses, have an error prone replication cycle which
15 results viable mutations in the genes which comprise the virus. This can result in alterations to antigenic determinants which have previously been used as vaccines. An alternative to the use of inactivated or attenuated pathogens is the identification of pathogen epitopes to which the immune system is particularly sensitive. In this regard many pathogenic toxins produced by pathogenic organisms during an
20 infection are particularly useful in the development of vaccines which protect the individual from a particular pathogenic organism.

The development of so-called subunit vaccines (vaccines in which the immunogen is a fragment or subunit of a protein or complex expressed by a particular pathogenic
25 organism) has been the focus of considerable medical research. The need to identify candidate molecules useful in the development of subunit vaccines is apparent not least because conventional chemotherapeutic approaches to the control of pathogenic organisms has more recently been stymied by the development of antibiotic resistance. A number of methods have been developed to identify potential antigenic
30 polypeptides which can be used as a vaccine. One such method is disclosed herein.

It has been known for many years that tumour cells produce a number of tumour cell specific antigens, some of which are presented at the tumour cell surface. The immune system recognises these antigens as foreign thereby resulting in the production of antibodies to self antigens, so called autoantibodies or autologous antisera.

One such technique is Serological identification of antigens by recombinant Expression Cloning, abbreviated to SEREX.

- Typically, the technique involves the extraction of RNA from tumour tissue followed by the selective enrichment of mRNA from the isolated total RNA. The mRNA is reverse transcribed into cDNA using viral reverse transcriptase. The cDNA thus synthesised is subcloned into an expression vector and transformed into an appropriate bacterial strain. The transformed bacteria are plated onto a suitable nutrient agar and under appropriate growth conditions the subcloned cDNA is expressed from the expression vector in the bacterial cell. The cells are lysed naturally by the use of phage based expression vectors, for example λ phage or phagemid based vectors, which through their lytic cycle cause cell lysis. The released polypeptides are transferred to a suitable membrane support (i.e. nitrocellulose, nylon) and exposed to autologous antisera from the patient from which the tumour tissue was originally isolated. The immunoscreening methodology allows the identification of genes that are over expressed or inappropriately expressed in a selected tumour tissue from a patient.
- We have exploited this technique to identify antigenic polypeptides expressed by pathogenic organisms during an infection. Autologous antisera produced during the infection is used to screen an expression library created from genomic DNA to identify and clone antigens.

In its broadest aspect the invention relates to the identification of antigenic polypeptides expressed during an infection by a pathogenic microbe and their use in vaccination.

5 According to a first aspect of the invention there is provided a method to identify opsonic antigens expressed by pathogenic organisms comprising:

- 10 (i) providing a nucleic acid library encoding genes or partial gene sequences of a pathogenic organism;
- (ii) transforming/transfecting said library into a host cell;
- (iii) providing conditions conducive to the expression of said transformed/transfected genes or partial gene sequences;
- 15 (iv) contacting the antigens expressed by the genes/partial gene sequences with autologous antisera derived from an animal infected with, or has been infected with, said pathogenic organism;
- (v) purifying the nucleic acid encoding the antigens or partial antigenic polypeptides binding to said autologous antisera; and
- 20 (vi) testing the opsonic activity of a polypeptide encoded by said DNA molecule.

In a preferred method of the invention said library comprises genomic DNA of a pathogenic organism.

25

Ideally said pathogenic organism is bacterial.

More preferably still said bacterial organism is selected from the following:

- Staphylococcus aureus*; *Staphylococcus epidermidis*; *Enterococcus faecalis*;
30 *Mycobacterium tuberculosis*; *Streptococcus group B*; *Streptococcus pneumoniae*;
Helicobacter pylori; *Neisseria gonorrhea*; *Streptococcus group A*; *Borrelia*

burgdorferi; *Coccidioides immitis*; *Histoplasma sapsulatum*; *Neisseria meningitidis* type B; *Shigella flexneri*; *Escherichia coli*; *Haemophilus influenzae*.

Preferably still said pathogenic organism is of the genus *Staphylococcus* spp. Ideally
5 organism is *Staphylococcus aureus* or *Staphylococcus epidermidis*.

In a further preferred embodiment of the invention said nucleic acid library is a lambda library, ideally a lambda expression library.

10 According to a second aspect of the invention there is provided a nucleic acid molecule comprising a DNA sequence selected from:

- (i) the DNA sequence as represented by the DNA sequences herein disclosed in Table 7 or Table 9;
- 15 (ii) DNA sequences which hybridise to the sequences identified in (i) above which encode a polypeptide expressed by a pathogenic organism and
- (iii) DNA sequences which are degenerate as a result of the genetic code to the
20 DNA sequences defined in (i) and (ii).

In a yet still further preferred embodiment of the invention said nucleic acid molecule is genomic DNA.

25

In a preferred embodiment of the invention there is provided an isolated nucleic acid molecule which anneals under stringent hybridisation conditions to the sequences herein disclosed.

30 Stringent hybridisation/washing conditions are well known in the art. For example, nucleic acid hybrids that are stable after washing in 0.1xSSC, 0.1% SDS at 60°C. It

is well known in the art that optimal hybridisation conditions can be calculated if the sequences of the nucleic acid is known. For example, hybridisation conditions can be determined by the GC content of the nucleic acid subject to hybridisation. Please see Sambrook *et al* (1989) Molecular Cloning; A Laboratory Approach. A common
5 formula for calculating the stringency conditions required to achieve hybridisation between nucleic acid molecules of a specified homology is:

$$T_m = 81.5^{\circ} \text{C} + 16.6 \log [\text{Na}^+] + 0.41 [\% \text{G} + \text{C}] - 0.63 (\% \text{formamide}).$$

10 According to a third aspect of the invention there is provided at least one polypeptide identified by the method according to the invention.

In a preferred embodiment of the invention, said polypeptide is associated with infective pathogenicity of an organism according to any previous aspect or
15 embodiment of the invention.

More preferably still said polypeptide is at least one, or part thereof, of the amino acid sequences represented in Tables 8 or Table 10.

20 In an alternative preferred embodiment of the invention said polypeptide carries a non-protein antigen, for example a polysaccharide antigen.

According to a fourth aspect of the invention there is provided a nucleic acid molecule characterised in that said nucleic acid molecule is part of a vector adapted
25 to facilitate recombinant expression of the polypeptide encoded by said nucleic acid molecule.

In a preferred embodiment of the invention said vector is an expression vector adapted for prokaryotic gene expression. Alternatively said expression vector is
30 adapted for eukaryotic gene expression.

Typically said adaptation includes, by example and not by way of limitation, the provision of transcription control sequences (promoter sequences) which mediate cell specific expression. These promoter sequences may be cell specific, inducible or constitutive.

5

Promoter is an art recognised term and, for the sake of clarity, includes the following features which are provided by example only, and not by way of limitation. Enhancer elements are *cis* acting nucleic acid sequences often found 5' to the transcription initiation site of a gene (enhancers can also be found 3' to a gene sequence or even located in intronic sequences and is therefore position independent). Enhancers function to increase the rate of transcription of the gene to which the enhancer is linked. Enhancer activity is responsive to *trans* acting transcription factors (polypeptides) which have been shown to bind specifically to enhancer elements. The binding/activity of transcription factors (please see Eukaryotic Transcription Factors, by David S Latchman, Academic Press Ltd, San Diego) is responsive to a number of environmental cues which include, by example and not by way of limitation, intermediary metabolites (eg glucose, lipids), environmental effectors (eg light, heat,).

20 Promoter elements also include so called TATA box and RNA polymerase initiation selection (RIS) sequences which function to select a site of transcription initiation. These sequences also bind polypeptides which function, *inter alia*, to facilitate transcription initiation selection by RNA polymerase.

25 Adaptations also include the provision of selectable markers and autonomous replication sequences which both facilitate the maintenance of said vector in either the eukaryotic cell or prokaryotic host. Vectors which are maintained autonomously are referred to as episomal vectors.

30 Adaptations which facilitate the expression of vector encoded genes include the provision of transcription termination/polyadenylation sequences. This also includes

the provision of internal ribosome entry sites (IRES) which function to maximise expression of vector encoded genes arranged in bicistronic or multi-cistronic expression cassettes.

- 5 These adaptations are well known in the art. There is a significant amount of published literature with respect to expression vector construction and recombinant DNA techniques in general. Please see, Sambrook et al (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbour Laboratory, Cold Spring Harbour, NY and references therein; Marston, F (1987) DNA Cloning Techniques: A Practical
10 Approach Vol III IRL Press, Oxford UK; DNA Cloning: F M Ausubel et al, Current Protocols in Molecular Biology, John Wiley & Sons, Inc.(1994).

According to yet a further aspect of the invention there is provided a method for the production of the polypeptides according to any previous aspect or embodiment of
15 the invention comprising:

- (i) providing a cell transformed/transfected with a vector according to the invention;
- (ii) growing said cell in conditions conducive to the manufacture of said polypeptides; and
- 20 (iii) purifying said polypeptide from said cell, or its growth environment.

In a preferred method of the invention said vector encodes, and thus said recombinant polypeptide is provided with, a secretion signal to facilitate purification of said polypeptide.
25

According to a fifth aspect of the invention there is provided a cell or cell-line transformed or transfected with the vector according to the invention.

In a preferred embodiment of the invention said cell is a prokaryotic cell.
30 Alternatively said cell is a eukaryotic cell selected from: fungal, insect, amphibian; mammalian; plant.

According to a yet further aspect of the invention there is provided a vaccine comprising at least one antigen or antigenic polypeptide according to the invention.

5 Ideally said vaccine further comprises a carrier and/or adjuvant.

The terms adjuvant and carrier are construed in the following manner. Some polypeptide or peptide antigens contain B-cell epitopes but no T cell epitopes. Immune responses can be greatly enhanced by the inclusion of a T cell epitope in the
10 polypeptide/peptide or by the conjugation of the polypeptide/peptide to an immunogenic carrier protein such as key hole limpet haemocyanin or tetanus toxoid which contain multiple T cell epitopes. The conjugate is taken up by antigen presenting cells, processed and presented by human leukocyte antigens (HLA's) class II molecules. This allows T cell help to be given by T cell's specific for carrier
15 derived epitopes to the B cell which is specific for the original antigenic polypeptide/peptide. This can lead to increase in antibody production, secretion and isotype switching.

An adjuvant is a substance or procedure which augments specific immune responses
20 to antigens by modulating the activity of immune cells. Examples of adjuvants include, by example only, agonsitic antibodies to co-stimulatory molecules, Freund's adjuvant, muramyl dipeptides, liposomes. An adjuvant is therefore an immunomodulator. A carrier is an immunogenic molecule which, when bound to a second molecule augments immune responses to the latter.

25

In yet a further aspect of the invention there is provided a method to immunise an animal against a pathogenic microbe comprising administering to said animal at least one polypeptide, or part thereof, according to the invention or the vaccine according to the invention.

30

In a preferred method of the invention said animal is human.

Preferably the vaccine, or antigenic polypeptide, can be delivered by direct injection either intravenously, intramuscularly, subcutaneously. Further still, the vaccine or antigenic polypeptide, may be taken orally.

Preferably the vaccine is against the bacterial species *Staphylococcus aureus*.

- 5 The vaccine may also be against the bacterial species *Staphylococcus epidermidis*.

It will also be apparent that vaccines or antigenic polypeptides are effective at preventing or alleviating conditions in animals other than humans, for example and not by way of limitation, family pets, livestock, horses.

- 10 According to a further aspect of the invention there is provided an antibody, or at least an effective binding part thereof, which binds at least one antigen or antigenic polypeptide according to the invention.

In a preferred embodiment of the invention said antibody is a polyclonal or monoclonal antibody wherein said antibody is specific to said polypeptide.

15

Alternatively, said antibody is a chimeric antibody produced by recombinant methods to contain the variable region of said antibody with an invariant or constant region of a human antibody.

- 20 In a further alternative embodiment of the invention, said antibody is humanised by recombinant methods to combine the complementarity determining regions of said antibody with both the constant (C) regions and the framework regions from the variable (V) regions of a human antibody.

- 25 Preferably said antibody is provided with a marker including a conventional label or tag, for example a radioactive and/or fluorescent and/or epitope label or tag.

Preferably said humanised monoclonal antibody to said polypeptide is produced as a fusion polypeptide in an expression vector suitably adapted for transfection or transformation of prokaryotic or eukaryotic cells.

Antibodies, also known as immunoglobulins, are protein molecules which have specificity for foreign molecules (antigens). Immunoglobulins (Ig) are a class of structurally related proteins consisting of two pairs of polypeptide chains, one pair of
5 light (L) (low molecular weight) chain (κ or λ), and one pair of heavy (H) chains (γ , α , μ , δ and ϵ), all four linked together by disulphide bonds. Both H and L chains have regions that contribute to the binding of antigen and that are highly variable from one Ig molecule to another. In addition, H and L chains contain regions that are non-variable or constant.

10

The L chains consist of two domains. The carboxy-terminal domain is essentially identical among L chains of a given type and is referred to as the "constant" (C) region. The amino terminal domain varies from L chain to L chain and contributes to the binding site of the antibody. Because of its variability, it is referred to as the
15 "variable" (V) region.

The H chains of Ig molecules are of several classes, α , μ , σ , α , and γ (of which there are several sub-classes). An assembled Ig molecule consisting of one or more units of two identical H and L chains, derives its name from the H chain that it possesses.
20 Thus, there are five Ig isotypes: IgA, IgM, IgD, IgE and IgG (with four sub-classes based on the differences in the H chains, i.e., IgG1, IgG2, IgG3 and IgG4). Further detail regarding antibody structure and their various functions can be found in, Using Antibodies: A laboratory manual, Cold Spring Harbour Laboratory Press.

25 Chimeric antibodies are recombinant antibodies in which all of the V-regions of a mouse or rat antibody are combined with human antibody C-regions. Humanised antibodies are recombinant hybrid antibodies which fuse the complementarity determining regions from a rodent antibody V-region with the framework regions from the human antibody V-regions. The C-regions from the human antibody are also
30 used. The complementarity determining regions (CDRs) are the regions within the N-terminal domain of both the heavy and light chain of the antibody to where the

majority of the variation of the V-region is restricted. These regions form loops at the surface of the antibody molecule. These loops provide the binding surface between the antibody and antigen.

- 5 Antibodies from non-human animals provoke an immune response to the foreign antibody and its removal from the circulation. Both chimeric and humanised antibodies have reduced antigenicity when injected to a human subject because there is a reduced amount of rodent (i.e. foreign) antibody within the recombinant hybrid antibody, while the human antibody regions do not illicit an immune response. This
10 results in a weaker immune response and a decrease in the clearance of the antibody. This is clearly desirable when using therapeutic antibodies in the treatment of human diseases. Humanised antibodies are designed to have less "foreign" antibody regions and are therefore thought to be less immunogenic than chimeric antibodies.

- 15 In a further preferred embodiment of the invention said antibodies are opsonic antibodies.

- Phagocytosis is mediated by macrophages and polymorphic leukocytes and involves the ingestion and digestion of micro-organisms, damaged or dead cells, cell debris,
20 insoluble particles and activated clotting factors. Opsonins are agents which facilitate the phagocytosis of the above foreign bodies. Opsonic antibodies are therefore antibodies which provide the same function. Examples of opsonins are the Fc portion of an antibody or compliment C3.

- 25 In another aspect of the invention there is provided a vector which is adapted for the expression of the humanised or chimeric antibodies according to the invention.

- In a yet further aspect of the invention, there is provided a cell or cell line which has been transformed or transfected with the vector encoding the humanised or chimeric
30 antibody according to the invention.

In a yet further aspect of the invention there is provided a method for the production of the humanised or chimeric antibody according to the invention comprising :

- 5 (i) providing a cell transformed or transfected with a vector which comprises a nucleic acid molecule encoding the humanised or chimeric antibody according to the invention;
- (ii) growing said cell in conditions conducive to the manufacture of said antibody; and
- (iii) purifying said antibody from said cell, or its growth environment.

10 In a yet further aspect of the invention there is provided a hybridoma cell line which produces a monoclonal antibody as hereinbefore described.

In a further aspect of the invention there is provided a method of producing monoclonal antibodies according to the invention using hybridoma cell lines
15 according to the invention.

In a further aspect of the invention there is provided a method for preparing a hybridoma cell-line producing monoclonal antibodies according to the invention comprising the steps of:

- 20 i) immunising an immunocompetent mammal with an immunogen comprising at least one polypeptide having the amino acid sequence as represented in Table 8 or 10, or fragments thereof;
- ii) fusing lymphocytes of the immunised immunocompetent mammal with myeloma cells to form hybridoma cells;
- 25 iii) screening monoclonal antibodies produced by the hybridoma cells of step (ii) for binding activity to the amino acid sequences of (i);
- iv) culturing the hybridoma cells to proliferate and/or to secrete said monoclonal antibody; and
- v) recovering the monoclonal antibody from the culture supernatant.

30

Preferably, the said immunocompetent mammal is a mouse. Alternatively, said immunocompetent mammal is a rat.

The production of monoclonal antibodies using hybridoma cells is well-known in the art. The methods used to produce monoclonal antibodies are disclosed by Kohler and Milstein in Nature 256, 495-497 (1975) and also by Donillard and Hoffman, "Basic Facts about Hybridomas" in Compendium of Immunology V.II ed. by Schwartz, 1981, which are incorporated by reference.

10 In a further aspect of the invention there is provided the use of the antibodies for manufacture of a medicament for the treatment of *Staphylococcus aureus*-associated septicaemia, food-poisoning or skin disorders.

15 In another aspect of the invention there is provided the use of the antibodies according to the invention for the manufacture of a medicament for the treatment of *Staphylococcus epidermidis*-associated septicaemia, peritonitis or endocarditis.

It will be apparent that the polypeptides identified by the method according to the invention will facilitate the production of therapeutic antibodies to a range of diseases resulting from pathogenic infection, for example, septicaemia; tuberculosis; bacteria-associated food poisoning; blood infections; peritonitis; endocarditis; sepsis; meningitis; pneumonia; stomach ulcers; gonorrhoea; strep throat; streptococcal-associated toxic shock; necrotizing fasciitis; impetigo; histoplasmosis; Lyme disease; gastro-enteritis; dysentery; shigellosis.

25

As has already been stated earlier, microbial organisms cause a wide variety of diseases. Listed below, and not by way of limitation, are a number of micro-organisms and some of the diseases they cause.

Micro-organism	Disease(s) caused
<i>Staphylococcus aureus</i>	Sepsis, food poisoning, septicaemia,
<i>Staphylococcus epidermidis</i>	Peritonitis, septicaemia, endocarditis,

	other hospital-associated diseases
<i>Enterococcus faecalis</i>	Endocarditis, cystitis, wound infections
<i>Mycobacterium tuberculosis</i>	Tuberculosis
<i>Streptococcus group B</i>	Sepsis, meningitis, pneumonia, bladder infections
<i>Streptococcus pneumoniae</i>	Pneumonia, meningitis
<i>Helicobacter pylori</i>	Stomach ulcers
<i>Neisseria gonorrhoea</i>	Gonorrhoea
<i>Streptococcus group A</i>	Strep throat, necrotizing fasciitis, impetigo, Strep. Toxic shock syndrome
<i>Borrelia burgdoferi</i>	Lyme disease
<i>Coccidiodes immitis</i>	Pneumonia
<i>Histoplasma sapsulatum</i>	Histoplasmosis, pneumonia
<i>Neisseria meningitidis type B</i>	Meningitis
<i>Shigella flexneri</i>	Gastro-enteritis, shigellosis, dysentery
<i>Escherichia coli</i>	Food-poisoning, gastro-enteritis
<i>Haemophilus influenzae</i>	Meningitis, pneumonia, arthritis, cellulitis

An embodiment of the invention will now be described by example only and with reference to the following materials, methods and tables:

- 5 Table 1 illustrates the immunization and bleed schedule for production of monoclonal antibodies reactive with peptide Hex A;

Table 2 illustrates an immunoassay of sera from mice immunized with peptide Hex A;

10

Table 3 illustrates an immunoassay of supernatants from anti-Hex A hybridoma supernatants;

Table 4 illustrates the immunization and bleed schedule for production of

15

monoclonal antibodies reactive with peptide 29kDa peptide;

Table 5 illustrates an immunoassay of day 98 sera from mice immunized with peptide 29kDa;

Table 6 illustrates an immunoassay of supernatants from anti-29kDa hybridomas supernatants from T75 Culture Flasks;

- 5 Table 7 represents the DNA sequences of *S.aureaus* partial gene sequences identified by the screening method;

Table 8 represents the protein sequences encoded by the DNA sequences illustrated in Table 7;

10

Table 9 represents the DNA sequences of *S.epidermidis* partial gene sequences identified by the screening method; and

- 15 Table 10 represents the protein sequences of the DNA sequences illustrated in Table 9.

Materials and Methods

20 Screening Genomic Libraries of *S. aureus* and *S.epidermidis*

- A λ ZAP Express library of genomic DNA of *S. aureus* 8325/4 and *S.epidermidis* was used. It contains fragments of 2-10kb from a partial *Sau*3A digest of total genomic DNA. This was cloned into the *Bam*H1 site of the vector. The library contains >10x coverage of the genome. The library was probed by plaque lift using an initial
- 25 screen of approximately 20,000 plaque forming units on a 9cm diameter Petri dish. The plating cells used, their treatment, the plating procedure and buffers were exactly as described in the manufacturers handbook (Stratagene). Plating cells, *Escherichia coli* XL1-Blue MRF', were infected with phage and plated in 3 ml top LB agar containing 10 mM MgSO₄ onto LB plates containing 10 mM MgSO₄. The plates
- 30 were then incubated at 42°C for 4 hr. An 8.5cm diameter nitrocellulose filter disc (previously soaked in 10 mM IPTG and air-dried) was placed on each plate and its location marked. The plates were then incubated for a further 3.5 hr at 37°C. The

5 filters were removed and washed in TBST buffer before blocking overnight at 4°C in TBST containing 6% w/v dried skimmed milk and 3% v/v pig serum (Sigma). The serum was used to block any Protein A clones on the filter. The filters are then treated with patient serum (1/5000 dilution) in blocking solution for 90 min at room temperature. Antisera have been obtained from patients convalescing from major *S. aureus* infections. The filters are then washed for 3x10 min in TBST. Secondary antibody used was goat anti-human whole IgG alkaline phosphatase linked (Sigma) at 1/30,000 dilution in blocking solution at room temperature for 30 min. The filters were then washed as above and developed using a standard colorimetric procedure.

10

Cross-reactive plaques were located on the agar plates and cored into 0.2ml phage buffer with 0.02 ml chloroform. The titre of each core stock was determined and the phage plated at approximately 200 plaques per plate. A plaque lift and screen was performed as above to give single, pure cross-reactive clones.

15

The pure clones were then spotted (1µl) onto plates to give a confluent plaque of 0.5cm diameter. 30 individual clones can be spotted on each plate. A plaque lift is performed and the filter probed with an appropriate sera. In this way clones can be tested for their cross-reactivity with other patient sera, non-infected donor sera and anti-Protein A sera.

20

Individual clones were then excised to give a phagemid in *E. coli* XL0LR using the manufacturers protocol (Stratagene). A plasmid miniprep of each was carried out and the size of the genomic insert determined by restriction mapping. The identity of the cloned insert was determined by DNA sequencing using primers against vector sequence, which allows sequencing across the insert. By comparison of the derived sequence against the public domain databases the nature of the cloned gene(s) can be determined.

25

30

Hybridisation Solutions/Conditions

Typically, hybridisation conditions uses 4 – 6 x SSPE (20x SSPE contains 175.3g NaCl, 88.2g NaH₂PO₄ H₂O and 7.4g EDTA dissolved to 1 litre and the pH adjusted to 7.4); 5-10x Denhardts solution (50x Denhardts solution contains 5g Ficoll (type 400, Pharmacia), 5g polyvinylpyrrolidone abd 5g bovine serum albumen; 100µg-1.0mg/ml sonicated salmon/herring DNA; 0.1-1.0% sodium dodecyl sulphate; optionally 40-60% deionised formamide. Hybridisation temperature will vary depending on the GC content of the nucleic acid target sequence but will typically be between 42⁰ - 65⁰ .

Mouse Model for Testing Candidate Vaccine Polypeptides

Mice are injected intravenously with 5 x 10⁷ *S. aureus* and mortality, bacteremia and abscess formation is monitored over the ensuing 7 days. At this dose 100% of the mice are bacteremic for greater than 4 days , 100% have detectable abscess formation in liver and kidney and greater than 80% of mice die within four days. At lower doses of injected organisms, bacteremia is detectable in the absence of death.

Immunization Program

Single proteins are injected at a dose of 10-100ug per mouse in RIBI adjuvant, boosted 14 and 28 days later and bled 14 and 28 days thereafter for evaluation of antibodies in their sera using ELISA. When groups of proteins are injected the final amount of each protein will be 10ug per mouse and the above immunization scheme will be followed.

Evaluation of Protective Efficacy of Single or Groups of Proteins

We will employ the mouse infection model described above to evaluate the protective efficacy of the proteins that are being tested. To this end groups of 5 mice will be immunized with single proteins or pools of 5 proteins as described above. We will monitor antibody titers to the injected proteins and when high titers are reached we will inoculate mice with *S aureus* at high and low dose. Control mice that have

not. been immunized or that were immunized with adjuvant only will also be inoculated with *S aureus*. We will measure levels of bacteremia, abscess formation and survival in all groups. All parameters of infection will be suppressed in mice that have high circulating levels of protective antibodies. If we find a pool of proteins that induces protection we will compare the protection induced by the individual components to that induced by the pool of proteins to see if protection was induced by a single protein or by the combined action of antibodies to multiple proteins. Using this approach we will identify protein epitopes that are protective.

10 In addition to using the *in vivo* model of mouse infection we will also obtain the sera from mice that are injected as above and monitor their sera for opsonophagocytic activity using a complement dependent system in the presence of human polymorphonuclear lymphocytes. This assay is well known in the art. This assay has been used an *in vitro* surrogate for measuring protective efficacy of antibody. Spleens
15 from mice that have opsonophagocytic antibodies will then be used as fusion partners in an attempt to make monoclonal antibodies that are reactive with *S. aureus*.

Using this multipronged approaches we will have a high level of confidence that we can identify protective epitopes that can be used either in a vaccine construct or that
20 can be used to generate monoclonal antibodies.

EXAMPLE 1

Immunoassay for detection of antibodies reactive with peptide Hex A

25 The binding of mouse sera or MAbs to Hex A was measured by immunoassay on wells coated with Hex A. One hundred microliters of a 250 – 500 ng/ml solution of Hex A in PBS was distributed into replicate Nunc Maxisorp Stripwells and incubated overnight at room temperature. The unbound material was removed from the wells by washing four times with PBS-T. Unbound antigen was removed from the plate by
30 washing four times with PBS-T. Antibody, diluted in PBS-T, was then added to the wells and incubated at room temperature for 30-60 minutes. After addition of the antibody, the wells were incubated at room temperature for 30-60 minutes in a draft-

free environment. The wells were again washed four times with PBS-T and ninety-five microliters of detection antibody was then added to each well. The detection antibody was either peroxidase-labeled goat anti-mouse IgG (gamma-specific), diluted 1:10000 in PBS-T, or peroxidase-labeled rabbit anti-mouse IgG₁, diluted 1:6000 in PBS-T.

Following another 30-60 minute incubation at room temperature, the wells were washed four times with PBS-T and each well received 100 µl of TMB substrate solution (BioFx #TMBW-0100-01). Plates were incubated in the dark at room temperature for 15 minutes and the binding reactions were stopped by the addition of 100 µl of TMB stop solution (BioFx #STPR-0100-01). The absorbance of each well was measured at 450 nm using a Molecular Devices Vmax plate reader.

Isotype was determined using a mouse immunoglobulin isotype kit obtained from Zymed Laboratories (Cat. No. 90-6550).

Immunization of Mice for Production of Monoclonal Antibodies Reactive with Peptide Hex A.

Five female BALB/c mice, approximately 8 weeks of age, were immunized with Hex A according to the schedule described in Table 1. All immunizations were administered subcutaneously in 50% RIBI adjuvant. Sera from the mice were tested by immunoassay, and based on the results of the assay described in Table 2, mouse 2021 was selected for hybridoma production. Mouse 2021 received a booster immunization of 32.5 ug of Hex A in PBS, administered intraperitoneally, three days prior to the production of hybridomas.

TABLE 1

**Immunization and Bleed Schedule for Production of
Monoclonal Antibodies Reactive with Peptide Hex A**

Experimental		Boost			
Day		(ug/mouse)	Adjuvant		Bleed
0		10 ug	RIBI		Yes
34		8.3	RIBI		Yes
48		None			Yes
60		25 ug	RIBI		Yes
74		None			Yes
98		25 ug	RIBI		Yes
124		None			Yes

TABLE 2

**Immunoassay of Sera from Mice
Immunized with Peptide Hex A**

Serum					
Dilution	2021	2022	2023	2024	2025
1000	3.553	3.569	3.226	3.336	3.439
3000	2.803	2.538	2.357	2.575	2.403
9000	1.663	1.336	1.314	1.522	1.357
27000	0.793	0.618	0.622	0.716	0.598
Buffer	0.095	0.078	0.145	0.066	0.089

Preparation of Hybridomas Reactive with Hex A Peptide

Hybridomas were prepared by the general methods of Shulman, Wilde and Kohler and Barta and Hirshaut (34, 48). Mouse 2021 was selected for hybridoma production based on the results of an immunoassay and received a booster immunization of 32.5 ug of antigen three days prior to sacrifice. Spleenocytes from

mouse 2028 were isolated and mixed with mouse myeloma cells SP2/0 (ATCC Catalog number CRL 1581) at a ratio of 10 spleenocytes:1 myeloma. The cells were pelleted by centrifugation (400 X g, 10 minutes at room temperature) and washed in serum free medium. The supernatant was removed to near-dryness and fusion of the
5 cell mixture was accomplished in a sterile 50 ml centrifuge conical by the addition of 1 ml of warm (37°C) polyethylene glycol (PEG; mw 1400; Boehringer Mannheim) over a period of 60-90 seconds. The PEG was diluted by slow addition of serum-free medium in successive volumes of 1, 2, 4, 8, 16 and 19 mls. The hybridoma cell suspension was gently resuspended into the medium and the cells pelleted by
10 centrifugation (500 X g, 10 minutes at room temperature). The supernatant was removed and the cells resuspended in medium RPMI 1640, supplemented with 15% heat-inactivated fetal bovine serum, 0.05 mM hypoxanthine and 16 µM thymidine (HT medium). One hundred µl of the hybridoma cells were planted into 952 wells of 96-well tissue culture plates. Eight wells (column 1 of plate A) received
15 approximately 2.5×10^4 SP/20 cells in 100 µl. The SP/20 cells served as a control for killing by the selection medium added 24 hours later:

Twenty four hours after preparation of the hybridomas, 100 µl of RPMI 1640, supplemented with 15% heat-inactivated fetal bovine serums, 0.1 mM hypoxanthine,
20 0.8 µM aminopterin and 32 µM thymidine (HAT medium) was added to each well. Ninety-six hours after the preparation of the hybridomas, the SP/20 cells in plate A, column 1 appeared to be dead, indicating that the HAT selection medium had successfully killed the unfused SP/20 cells.

25 Ten days after the preparation of the hybridomas, supernatants from all wells were tested by ELISA for the presence of antibodies reactive with peptide Hex A. Based on the results of this preliminary assay, cells from three wells were transferred to a 24-well culture dish and expanded. Supernatants from these cultures were retested by ELISA for the presence of antibodies that bind to peptide Hex A.

30

Using IgG-1-specific detection, the absorbance values obtained with the supernatants from hybridoma culture 02-101FE1, 02-101ED8 and 02-100JC10 were 2.150, 2.230 and 2.574, respectively, compared to an absorbance of 0.044 with buffer alone (Table 3). Absorbances were lower, but still positive, with gamma-specific detection (Table 3). Each of the cultures was expanded, cryopreserved and cloned by limiting dilution. Two-three clones of each culture were expanded and cryopreserved for future evaluation.

TABLE 3**Immunoassay of Supernatants from Anti-Hex A Hybridoma Supernatants**

		Detection With	Detection With
Culture ID	Dilution	Anti-Mouse IgG-1	Anti-Mouse Gamma
02-101FE1	2	2.150	0.941
02-101JC10	2	2.574	1.403
02-101ED8	2	2.238	1.174
Buffer		0.044	0.073

EXAMPLE 2**Immunoassay for detection of antibodies reactive with peptide 29kDa**

The binding of mouse sera or MAbs to 29kDa was measured by immunoassay on wells coated with 29kDa. One hundred microliters of a 500 - 1000 ng/ml solution of 29kDa in PBS was distributed into replicate Nunc Maxisorp Stripwells and incubated overnight at room temperature. The unbound material was removed from the wells by washing four times with PBS-T. Unbound antigen was removed from the plate by washing four times with PBS-T. Antibody, diluted in PBS-T, was then added to the wells and incubated at room temperature for 30-60 minutes. After addition of the antibody, the wells were incubated at room temperature for 30-60 minutes in a draft-

free environment. The wells were again washed four times with PBS-T and ninety-five microliters of detection antibody was then added to each well. The detection antibody was either peroxidase-labeled goat anti-mouse IgG (gamma-specific), diluted 1:10000 in PBS-T, or peroxidase-labeled rabbit anti-mouse IgG₁, diluted
5 1:6000 in PBS-T.

Following another 30-60 minute incubation at room temperature, the wells were washed four times with PBS-T and each well received 100 µl of TMB substrate solution (BioFx #TMBW-0100-01). Plates were incubated in the dark at room temperature for 15 minutes and the binding reactions were stopped by the addition of
10 100 µl of TMB stop solution (BioFx #STPR-0100-01). The absorbance of each well was measured at 450 nm using a Molecular Devices Vmax plate reader.

Isotype was determined using a mouse immunoglobulin isotype kit obtained from Zymed Laboratories (Cat. No. 90-6550).
15

Immunoassay for detection of antibodies reactive with peptide 29kDa

The binding of mouse sera or MAbs to 29kDa was measured by immunoassay on wells coated with 29kDa. One hundred microliters of a 500 - 1000 ng/ml solution of 29kDa in PBS was distributed into replicate Nunc Maxisorp Stripwells and incubated
20 overnight at room temperature. The unbound material was removed from the wells by washing four times with PBS-T. Unbound antigen was removed from the plate by washing four times with PBS-T. Antibody, diluted in PBS-T, was then added to the wells and incubated at room temperature for 30-60 minutes. After addition of the antibody, the wells were incubated at room temperature for 30-60 minutes in a draft-
25 free environment. The wells were again washed four times with PBS-T and ninety-five microliters of detection antibody was then added to each well. The detection antibody was either peroxidase-labeled goat anti-mouse IgG (gamma-specific), diluted 1:10000 in PBS-T, or peroxidase-labeled rabbit anti-mouse IgG₁, diluted 1:6000 in PBS-T.

Following another 30-60 minute incubation at room temperature, the wells were washed four times with PBS-T and each well received 100 µl of TMB substrate solution (BioFx #TMBW-0100-01). Plates were incubated in the dark at room temperature for 15 minutes and the binding reactions were stopped by the addition of 100 µl of TMB stop solution (BioFx #STPR-0100-01). The absorbance of each well was measured at 450 nm using a Molecular Devices Vmax plate reader.

Isotype was determined using a mouse immunoglobulin isotype kit obtained from Zymed Laboratories (Cat. No. 90-6550).

Immunization of Mice for Production of Monoclonal Antibodies Reactive with Peptide 29kDa

Five female BALB/c mice, approximately 8 weeks of age, were immunized with 29kDa according to the schedule described in Table 1. All immunizations were administered subcutaneously in 50% RIBI adjuvant. Sera from the mice were tested by immunoassay, and based on the results of the assay described in Table 2, mouse 2028 was selected for hybridoma production. Mouse 2028 received a booster immunization of 50 ug of 29kDa in PBS, administered intraperitoneally, three days prior to the production of hybridomas.

TABLE 4

Immunization and Bleed Schedule for Production of Monoclonal Antibodies Reactive with Peptide 29kDa

Experimental		Boost			
Day		(ug/mouse)	Adjuvant		Bleed
0		10 ug	RIBI		Yes
34		10 ug	RIBI		Yes
48		None			Yes
60		20 ug	RIBI		Yes
74		None			Yes
98		20 ug	RIBI		Yes

TABLE 5

Immunoassay of Day 98 Sera from Mice

Immunized with Peptide 29kDa

Mouse ID		Sera at 1:1000		Sera at 1:10000	
2026		0.260		0.078	
2027		1.415		0.306	
2028		2.184		0.383	
2029		0.838		0.107	
2030		1.073		0.154	
Buffer		0.061			

Preparation of Hybridomas Reactive with 29kDa Peptide

Hybridomas were prepared by the general methods of Shulman, Wilde and Kohler and Bartal and Hirshaut (34, 48). Mouse 2028 was selected for hybridoma production based on the results of an immunoassay and received a booster immunization of 50 ug of antigen three days prior to sacrifice. Spleenocytes from mouse 2028 were isolated and mixed with mouse myeloma cells P3X63Ag8.653 (ATCC Catalog number CRL 1580) at a ratio of 10 spleenocytes:1 myeloma. The cells were pelleted by centrifugation (400 X g, 10 minutes at room temperature) and washed in serum free medium. The supernatant was removed to near-dryness and fusion of the cell mixture was accomplished in a sterile 50 ml centrifuge conical by the addition of 1 ml of warm (37°C) polyethylene glycol (PEG; mw 1400; Boehringer Mannheim) over a period of 60-90 seconds. The PEG was diluted by slow addition of serum-free medium in successive volumes of 1, 2, 4, 8, 16 and 19 mls. The hybridoma cell suspension was gently resuspended into the medium and the cells pelleted by centrifugation (500 X g, 10 minutes at room temperature). The supernatant was removed and the cells resuspended in medium RPMI 1640, supplemented with 15% heat-inactivated fetal bovine serum, 0.05 mM hypoxanthine and 16 μ M thymidine (HT medium). One hundred μ l of the hybridoma cells were

planted into 952 wells of 96-well tissue culture plates. Eight wells (column 1 of plate A) received approximately 2.5×10^4 P3X63Ag8.653 cells in 100 μ l. The P3X63Ag8.653 cells served as a control for killing by the selection medium added 24 hours later.

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Twenty four hours after preparation of the hybridomas, 100 μ l of RPMI 1640, supplemented with 15% heat-inactivated fetal bovine serums, 0.1 mM hypoxanthine, 0.8 μ M aminopterin and 32 μ M thymidine (HAT medium) was added to each well.

Ninety-six hours after the preparation of the hybridomas, the P3X63Ag8.653 cells in plate A, column 1 appeared to be dead, indicating that the HAT selection medium had successfully killed the unfused P3X63Ag8.653 cells.

Ten days after the preparation of the hybridomas, supernatants from all wells were tested by ELISA for the presence of antibodies reactive with peptide 29kDa.. Based on the results of this preliminary assay, cells from 3 wells were transferred to a 24-well culture dish and expanded. Several days later, supernatants from these cultures were retested by ELISA for the presence of antibodies that bind to peptide 29kDa.

The absorbance values obtained with the supernatants from hybridoma cultures 02-100EC7, 02-100HH10 and 02-100FG5 are presented in Table 3. Based on these results, cultures 02-100EC7 and HH10 were expanded, cryopreserved and cloned by limiting dilution. Two-three clones of each culture were expanded and cryopreserved for future evaluation.

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TABLE 6

Immunoassay of Supernatants from Anti-29kDa Hybridomas

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Supernatants from T75 Culture Flasks

Culture ID	Culture Dilution	Detection With Anti-Mouse IgG-1	Detection With Anti-Mouse Gamma
02-100HH10	2	1.021	0.312
02-100EC7	2	0.687	0.230
02-100FG5	2	0.048	0.048
Buffer Alone		0.044	0.050

TABLE 7

LOCUS 1 (E8/B1/I16)

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LOCUS 3
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LOCUS 4 (E103)
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LOCUS 5 (L4)
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LOCUS 6 (D1)
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LOCUS 7 (D3)

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LOCUS 8 (D4)
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LOCUS 9A (D22)
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LOCUS 9B (I2)
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LOCUS 9C (J13)
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LOCUS 9D (M11)
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LOCUS 9E (M13)
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LOCUS 10 (D9)
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LOCUS 11 (D10)
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LOCUS 12 ()
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CACTAACCTTAGCTTTGATTGATATAAATCATCCGATAAATCTGTATATAGTTTTTTAAT
TTGGTCTGATCGTAACCAACCTTTATATACAATTGTTTTGCGTGATAAGCTCGTAAATA
CAATTCTAAATCGCACTGAGTCGAATAGAACTCTAATTGTTTTCTCGCTAAAAACAAACG
CTTTTCAACATCTTCAATGTCCCTAATATCAATAAACACTTGTTGAATGACTGGCATCGT
ATCTGCTACATGTTTAGCAATGGCATCTTTATTAACCTGGTACATTACGATAACCAAGAAT
TGATAACCCTTCGCCCTTCAAAATATTTTTTAAAACTACTTCATGTTTCAAGAACCTAAAT
GCGTTCTTTGGAAAAAATAACCCACGGCATATTCACCTTCACCTGGGATATCAAAGTC
CGTTACATGTTGTTTGAAAAATGCAAAAGGTATTTTCAGTCATAATACCTGCGCCATCACC
AGTGATGCCATCTGCGCCGACCCCGCCCT

LOCUS 13 (D18)
GATCCATTGTTTCGCAGCAGCTGATGTCATTTTCATACATAACTTGTGAAATACCATGAAAA
GACGGATTTCGTTATACTTTCACTTGCTCCAGGAATCATAAAAGCAAGTGCTGAAAATACT
AAAATTAAAATTGGGTGTATGAGAAAGACTAAGACAATACATTTTCATTTACGGGCGCCA
ATTGGCATATTTAAATATTCTGGTGTTTTACCAACCATCAAACCTGCATATAAACACCGTC
AGTAAGACAAATATCAATAAATTCATGAGTCCTACGCCTTCGCCACCAAATACAACATTT
AGCATCATTAATACCATTTGGTCCTAATCCACCTATAGGCGTTAAGCTATCATGCATGTTA
TTAACAGAACCCGTTGTAAATGCCGTCGTAATAACTGTAAATAGTGCTGACAAACCTGCT
CCAAACCGTACCTCTTTACCTTCCATATTCGGTCCATAAATGCCTAAATTCGCTAGTATT
GGATTACCACGATACTCACTCCACATAGTTAATGTAAGAATTGCTATAAAAAATGAAAAAC
ATTGCGACAAATAATATCAACGCATGACGATGTACTCGTTTACCATGTCTACTTAACATG
CGACCAAATAAGAACAAACATTGACATAGGAAGTAACATCATACTGCCCATTCTCTATAAAA
TTGCTCCAAATATTTGGATTTTCAAAGGTGTTGCAGAATTTCTTGCTAAAAATCCTCCA
CCATTTCGTACCAAGATGTTTTATTGATTCAAGTGATGCAATAGGTCCAAATGCAATATGT
TGAATATGTCCGCTTAAAGTCCGAATCATTAATTAGCATGCAACGTTTGTGGTACACCT
TGAGTCATCAATAAAATACTAATTAAACATGATAATGGTAAAAGTACTCGGACAATAAAC
CGAACAAATATCTTGATAAAAATTACCAATGATATTAGTTAATCCAGTTAAACGTCTCAAC
ATCGCTATACAAACGGCGTAACCTGATGCACTAGATGTAAACATTAAATATGTCATTACA
ATCATTTCGCTTAAATATGTCACATCTGATTCACCGTTATAGTGTTGTAAATTACTATTT
GTTAAAAAAGATATTGCTGTATTAAACGCTAAATCTATCGATTGGTTTAAATTATGATTT
GGATTTAAAAAAGCCATTGCTGAACTATTAGCAATACAAATGTTATAAACCCCATAAAT
CCATTAAATGCCAGAAAATGTTTGACATATGTTTTAGCTGACATGTGTTCTAAATCTGTG
CCGATAATTTTAAACACATATTTTCAAATCTAGTAAATATTAAATCTACTCTTGACGAT
TGCACCAATGCTACGCGATATAGATATCCACTAAAAACATACGTAATCATAACCATCATT
GTTAGAAACAAATTATTTCCATGATAACCCTCACTTAATATATTTCTAAAATTTTTCAC
TACGAATTAAGGCATAAAATAAATACAAAATAATGCAATAACTACCAGTAATAAACGA
TGAGCATTGCCATAACCTCCTTACAACACAACATCGTAACAACCTTGTTTATGAGAGA
AATATTAATTTTCAAACCTAGTTATTAAGAAATCATTAAGATGTGTATGCAGAAATAAAT
TTTATAGCATTTAATTGTGAAGAATATTATGATATTGCTATCGAGGTGAAGGTTATGTCA
AACACTGAATCGCTAAACATAGGAAAAAAGCGTGGATC
LOCUS 14 (D21)
GATCACTGCATCTCCATCATTAACACCGTCATTTTGATTCTCAACGATGAATGGTACTAC
GAATTCGTCAGTTAAGCCCTCATTATAGCTTGCTTCTACACCTTCTTTGGCAGTTGCATA
AGTTGGGGCATCAAAATTACGAATAGCATTGTAAGCTTTTTCTTCACGTTCCCAACGTTT
GTCACGATCCATTGCATAATAACGACCAGACACAGATGCAAATTGACCAATGCCTAATTC
ATTGAATTTAGCTTCAGTCTCTTCGATGTATTTCAAAGCGGATTTTTGATCTACGTCACG
GCCATCTAAAAATGCGTGTACGTAACTTTTTCAACACCTTGTTTTTTAGCAAGTTCTAA
CAAAGCAAATAAATGTTTGTAATGACTGTGTACACCACCGTCAGACAATAAACCAAAGAT
GTGTAACGCTGAATCATGTGAATTCACGTGTGCAATTGCATTATTTAAACATCATTTTC
AAAGAAATCACCGTCTTCAATTGATTTATTGATTTCGAGTTAAACTTTGATAAACGATACG
TCCTGCACCGATATTCATATGACCAACTTCTGAGTTACCCATTTGTCCTTCAGGTAGTCC
AACATCTAAGCCACTCGCTTCGATTTGAGTCGTTGGATATTTGTTGTAATAACGATCAAA
ATTAGGCTTGTTTGCTAATTTTACCGCATTACCATGTTTCGCTTTTCGCGGTTTCGCAAAACC
ATCTAAATAAATTAACGCAGTTGGTTTCTTAGCCATGATTATTTTGACACCTTCTAACAAT
TGTACGAAATCTTCAACTTTAAGTGATGCGCCACCTACTAATGCCCCATCAATATCAGTT
TGTGCCATGTATTCTTTAATGTTGTTAGGTTTAACACTACCACCATATTGAATACGAGTT
GCTTCTGATACTTCTTTGCTTGATAAGTCAGCAATAGTTTGACGTACAAATGCACACATT
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GCAATTACAACCTGATTTAAGTTGATCTTCAGATAAACCTGCAACAGCTTTCTTAACTTGC
 TCACCTACAACATCGTTAGCTTTACCACTTTCACGCTCTTCGTCTGTTTCACCAACACAT
 ATAATTGGAGTCATTCCATGTTTGAAAATAGCGTGCGCTTTTTTGTAAATTTCTTCATCT
 GTTTCGTGGAATAATTCACGACGTTTCAGAATGACCGATAACAACGTATTTAACGCCTAAA
 TCTGCTAATGCAACTGGAGACGTTTCACCTGTGAACGCACCATTATCTTCGAAATACGTA
 TTTTGAGCACCGATTTCTAAACCTTGTGCTTTTCCTTCTTTAACTGCAGTAGTTAATGCA
 TCTAATTGAATTGCTGGTGCACAAATTACTGATTCTACTTCTTTTGAATCTGGTAGTGTT
 GGTAATGTATTGACGAAGTCTTTTGCTTCTTGTACTGTTTTGTTCATTTTCCAGTTACCA
 GCTATAATTGGTGTCTCATTAAAGACACTCCTTGTTTTGTAATATTTTGAAGTGA
 TGAAACACGATGTCATCTTGTGACTGTTTTCCCGTAACAATGTTAAACAAACATGCCACA
 TCACTTTAAACTATCACTTTATTATTATTATTGATTGCTTTGATACCAGGCAATTCTTT
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 AAAACCTAAAGAGATTGCTGCTGCAGCTGAATCACCGCCACCGATAATCGTAATTGCATC
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 TAATTTTACAGTGTTTGGTCCAATATCCATACCTTCTTGGTCTGCTGGAATTGAATCAGA
 TGGTACTACAGTGATTTTGGCATCATTAGAAAATTCTTTAGCAACTTTAGTGTCTACTGG
 TAATACAATTTTATCACCATGTTTTTCTAATAAATCTTTTGCGAAGTCGATTTTATCTTC
 TTCTAATAATGAAATACCAATTTCTTTACCTTGCCTTTTAAAGAAAGTATAAGCCATACC
 TCCGCCGATGATAATTTTATCAGCTATGTTAACTAAGTTTTTGATGACATTAATTTTGTC
 AGATACTTTTGCTCCACCTAAATAGCAACAACCTGGTTTATGTGGATCGTTAACTACGCC
 GCCAATAAACTTAATTTCTTTATCCATTAAGAATCCAGCTGCAGTTTCTAAATGTGTAGA
 AATACCAACATTAGATGCATGCTCACGATGCGCAGTACCAAAGCATCATTTACAAACAC
 ATCACCTAAAGATGCCCAGTATTTACCTAATTCTGGATC

LOCUS 15 (I1)

GATCCTGAAACGTAATTAATTGAACTGTAGAACCTTCAGTCACCTTGTGTCTTTTCTA
 ATCACTACTACTGGTAAATTTAAATATTAGCAACCGCATTTGCCAATGAAATACCTTTT
 GTCGCAATGGTAACAACAGCATCTAATTTTTCTTCCATGTAAATACTGGCAATTAACCTA
 CCAACTTTGTTTAGTAGCGATGGATTACCTACCAAATCTGATAAAAATAAATATCCGCCA
 GGTAACAAACGTTCTTTCTTCTAATAGAGTAATGACCTCATTAACAACCTTCAGTCGCC
 TCTTCTTTACTCATCATTTGGTTTATACGTAACACCACCACCTTGCGCCAGCAGTAGTAATT
 ACTGTACCTAACTTTTCTTTTGGGAATGTATTTTTTATAATTTGGACATCTTCACTTATT
 GAAGACTTCGCCTGTTTAAATTTTTTCACAAAAAAGTTAATGGAATCAATTTATTGCGA
 TGGTTCATCAAATATTGCGTCATAAAAACAATTCTCTCGCTTCGTTTATATCTCATCTTT
 TCAACCCTTCTATCCTAATAGTCTAATAAGTACACTTCATTACAACAACCGTTAACTGC
 ATTATAAATATTTTTTGCTTGGCTTTCTTTTCGTGCTAGCCCATACACAGTAGGTCCGCT
 TCCACTCATTAACGCACCATCTGCACCACTTTTCAACATATTATTTTTTAATTTATCGAT
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 TTGATAATCTCGATTTTCTAAGGCCTCATAACACATTTTCGTATGTACGTCGTAACGCTT
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 TGCTGCATCAGCCGAACCTCCAGCTAAGCCAGCAGAAACAGGTATTTCTTTATCGATAGA
 AATTGTTACACCTTGCTTTAGTTGATATTGCTCAATAAATAGTTGCGCTGCACGATATGC
 GAGATTTTTATGATTAGAAGGCACATAATTATGTTCAATCTCAACAACCTATCTTTGATC
 TTTTCTTTTATGAAAAGTTAAACGATC

LOCUS 17 (I3)

GATCGACAACACTCTAAATATATAGAAAATAGGTATTAATTTAACTATAAATCTAAATAA

TAATGCAAAGATGATTAAAATAACGATAGCTAAAGCAATACCAATAATAAAATCTTTGGT
CGCTAGCTCACCTATCATCCCCATATAGAAAATGATAACCTCGACACCTTCACGCAACAC
AGATATTAAACCAATCGTCGCTAACAATACCAAATTACCATTACTAATCGCATTAGCATA
CATATTTTAAATCATGTTCATTCCAACGTTTTGTCATTGTAACGTTTGTGCATCCAAACACC
AACGATAAACATTAATATGACCGCAACGATACCTAATCCCGCTTCCATACTTTCACGAAG
AATGCCACTATTCCCTAAAGTTTCTACAAACGTAATTGCTAAGATAATACTCAGTACAAG
TCCGGCAATTGCACCACCAATCACACTTGCAGTCCCTTTCTTATCTTTTACATTACGCGT
CATGGTAGTCAATGTTCATTACAATTAACAACACTTCTAGCCCTTCACGTAAAAAGATAAT
CATCACATCGACGAAGCTATAACTATGGCCAACAACCTCTTTAATTTGGTTATTTAAATC
TACTAAACCATCTTTCACATGTGCTTTATTATGTTTCGTCTAATACACTTTGATAATATGG
TATTTTATCTTCAATTTTCGTATACAAAGCACCGTCTTTAGTTTGAATTTGACCTTCAAC
ATACGGCCAAGTTTCTATAAAATGTGTAAGCGCAGCATCAGCATCCGACAATTGATTGTC
GTCGATAGCTTTAATCGCCTTCTCTAACGCATCATTTAATTGTGATACATGGTATTGATC
ATTTGCAGACGTATTACTTTTTTTATCGACATGATCAATATTTGATTTAAAAGTTGTCCA
AGCATGTGACACTTTTGCCGTATCTAATGGTGACTTATGAATTGCAATTCTAAGTTGTAA
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CACTTGCTCAATTGCTTTCTGCTTATTGTCATTTCGATATCGAATTATTAGAAAGTGCAGA
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CAGAACCTCTATGTGTAATATATTCAATTTAATTTATCATTACTACCTAAATTATTTTGTA
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TACCTGTGCGGTCATCCGTACCATCCACATAATTAAAGGCTCTACGTAAAATTGACGTAT
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CTTTCGCTTTTAAGTCAATTTTCATCAAACCTTTCCACCTGTTAACGGTGCACCACTAT
GTCGTTTCCGACCAAATGTAGCCTCTTGTTCTTCCAGCGCAGTACGATC
LOCUS 18 (15)
GATCGTTTAAATGTTCAATATATTCCGCTGCACTTTGCGCTGCAATACTACCATCGCCAG
TAGCAGTGACAATTTGGCGTAAACCTTTGTGCGGAACATCTCCTGCTGCAAAAATACCTG
GTAATGATGTTGTCATATCATCTTTTGTGTTACAATATAACCAACATCATTTGTAATACCTA
AGTCTTTAAATGGCGCTGTTAATGGTTTCATACCAATATAGATGAATACACCATCAGCCT
CGTGTGTTTCTTCTGAACCATCTTTTGTAGACGTTAATGTCACAGAACCCACTTTGCCGT
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CTTTGTCAGCAAATTTAGTTAAGAATGTTCCCTCTTCTACTGCTGAATCACCACCACCGA
TAACGAATAGGCGTTTATTTTTAAAGAATGCACCATCACATACTGCACAATAACTTACAC
CGCGTCCACCAAGTTCTTGTTCAACCGGAACACCAATTTTCTTGATTCTGCACCTGTAG
CAATAATAACCGCTTTCGCTGTTAATTCTTTATTACCAAAGTTAATCACTTTATATTGCG
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CTACTTCTTCTGTATTAGCCATTTGACCGCCTGGAATACCTCTTTCAATCATAACTGTTT
TTAAATTAGCACGTGATGCGTATACTGCAGCAGTCATACCAGCTGGACCTGCACCGATAA
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GCGCATTATATAATAAATCTAACTTTTCATAAATCTATATGCTCAAGAGAAATTCAATCA
TTTTGTTGAGTTTATATTGTGTTATGCCTAACCATGTTGTAATTTGCTTCTTTGTAACGT
TTTCGAGGTTGATTTTAAAATACAAATAAATAAACGCACCGATATATGGCTCAACATCAG

TTAAATCTACTTTTTTCAGCAATTATGAGTTCACCTTGATTAATCCATGCAACCATTACAT
CATTTTCACTTACAAATAATTCAATTATGGTAAAGCGTTAATAAGCCGCGATGAATGAAGT
CTAATTTATTGAGCGTTAAACCTTGAACATAACGTTAAATACAATTTCTCATAATTAT
TTAGATTTTCCAAAACCTGCCAAATACTTTCCGTCATCACAATTTCTTTACCATTTAATT
GATC
LOCUS 19 (I8)
GATCGTTGATTTGATTAGTGATGGTTGAACAAATTAAAAATAAACTACTTACTGCAAATA
CTACGCCCATACGATAAACGTTAGTAGCTGGTGTAGTATAACTTTGTAATGGCAGCGCCAC
TAAGACTGCCAATAATTTGACCAACAATAACATACTGTTCTGTCGTTCCAACAAATGTGC
CTTTAAGTTGTTGATGACACGCATTCACGACAACAAACATGACACTTTGAATCAATGCAC
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CGGACTGTAAAAATCCAATCACACTACGGTCATCTATCGCTGTATGATTCATGATGAAG
CAAGTGGTGATAATGCAGTTAGCATGCCATACATAGCAAAGTTTGCTAAAACGCCAACGA
TAATAAATCGACATGTTTGTGTTGTGCATAATAGACATTGAAATGAACGGCGAATACCTT
TATTAATATTTGGTGTGTTGTGATTTTGGCATATGTGTGCTTTCAATCAATTTTAATGCAC
CGAAAATACAGACAATAAAAGTAATAACGGCAATACTCATCAGTAACGCACTAAACCTA
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CAGCAATCGCTATAAACTGACTGAGCCATAAAATGCGAAAGTTACTGCGCCATATAGACT
GATTAATCATATGTCACCATTGGATTGTTGGTACGGTAGTTAAACCTGAAGGCATACTACCT
CCACCCTATCACGTTGATATAGCAATGGTAATAAAATTTGTTTGAATGGCCACGTCTGT
TTATCAAATAAAATGTGTCTGACAGCTAGCTGATCAGTTGTAACCCAGGAAATAGTTGCC
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AAATCTTGAATTGCATCAATAATGGCATATAGATTTACCGATACAGCTAATGTTTGAAA
TAAGCAAAGAATGTTTCCAAATCCTCATTAATTAGCGTATTAGGTGTATCTTCTCTGACG
ACATACTTCGGCAATGAAAGCTGATGTGCTGTTAGCCATGGTTTATAAATTCTGACAGTA
TCATGATCACGTAACACGCATTTTGTACACGTCCATCTTCAAATGACAACAATATATTT
TGACCATGCAACTCTGGTAATGCGCCGTATTGCATAAATGATAGTGTTACCTTTAAAAAG
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CAAATCATTTGATATAAAGTGCGATCATTTGCCGCGAGTGCTGCCATTGACACTAGCTGT
TGCGTATCATTTTTGGCTAGCACTTCGGGATACTTTCTTAGCTGAACAGTTAGATGACCT
AATTGATCTTTGAAAATATCATTATCTTGACCCATATATGACCACCAAGCTGTTTCATCA
CAAACCATGACATACTTAGCTAGTGCTTCATCTTTTTCTATAAGCTGACGTAATAATTGT
TCTGCTTGTCTCCGTTTTTTCATGTAACGCGTAGGCGTTAGCCTTAATGCGCCTAATGAC
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TCTTCAAATAATAATGCATCAACTAAATCTCTTAATATTATCGCTTGTGCTGTATTGACT
GCTGTATGATTCTGCAATGTTTCAGACACCTCGCATTCTTAATATAGGTTCAATGTTGTCC
CAATATTTTGTGTTGTGCTGTTGATAAATAAAATAAGCACTTGAAATATCTTCGATAG
CCATACCCATCGGATTAAGTAATATGATC
LOCUS 20 (J7/M10)
GATCGCTTACAAAACATAACAAGCTTTAAAGATATTGCCAAAATTCTTTATTCAACGAGA
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CGGGACTATTTTTATGGGAGACATTACGTAATAATTTCTATTACTCTGCTATCAATGTAC
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AAGACGAACTTGGAGACTTACCACAATGGATTAGTGATTTAGATGGTGGCTTTTATAAAC
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TGGAAATGACCATTACACAAGTATGGTTATTTATGCAGATGGTAACAATTTCAAGTGTGG
GTGCTAACCTTTTCTTAATGAAAAAGGCGCATGAAGACGGTCTTGTAGATGATGTCGTTG
CACAATCAATTGATAAATTACATTATAGCTTTAATCGTTTGAAGTATAGTTTGAAACCAG
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TCAATACAGCACAACGTGTGCGCTCAAACGTGCGAAATATGAAGCAGAAACAA
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ATCAACGTTACATTCAATCGTTGGAGAAAATTGGCTTTATTGACTTACTAAAATCTAAAA
AATCATATGAAAGAATTGCACATATGTTAAAACTGGTAAGCCATTACGTAATTAAAAGA
TAGTCATTAAGAGAGGATGATAACCATGCAAGAAGCATACATTGTAGCTTATGGGCGTTC
AGCCGCAGCGAAAGCAAAGCAAGCGCATTATTCCACGAAAGACCTGATGATGTCGCAGC
CAAAGTATTACAAGGCGTATTGAAACGTATTGACGGAAAATTCAATAAGAATATGATTGA
AGATGTCATTGTTGGTACGGCTTTTCCAGAAGGATTACAAGGCCAAAACATTGCACGAAC
GATTGCATTGCGTGCGGGATTATCTGACACGGTACCGGGTCAAACAGTGAATCGCTACTG
CTCATCAGGATTACAAACCATCGCGATTGCAGCCAATCAAATTATGGCTGGTCAAGGAGA
TATACTTGTAGCTGGTGGCGTTGAATTGATGAGTGCCGTACCAATGGGTGGCAACGAGCC
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TGCTGAAAATGTAGCATCCCAATTTGACGTATCACGCGAAGATCAAGATGCTTATGCTGT
CAGAAGTCATCAACGTGCCTATGACGCACAACGTGATGGTCGGTTCAAAGATGAAATTAT
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LOCUS 21 (G3)
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LOCUS 22 (119)
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LOCUS 24 (L10)
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LOCUS25 (HA4)
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LOCUS 26 (L19) :
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LOCUS 27D (AF7)
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LOCUS 28 (H130)
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LOCUS 29 (A) N10
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LOCUS 29 (B) GE2
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LOCUS 30 (N15)
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LOCUS 31
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LOCUS 32A (HE9)

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LOCUS 32B (P9)

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LOCUS 33 (014)
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LOCUS 34 (O18)
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LOCUS 35B (P15)
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LOCUS 36 (P5)

GATCATCTCTATCAATTTTTATATTAAATTCATTTTTTTGAATCGATAAAATAAACTCGA
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LOCUS 37 (P8)
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LOCUS 38 (P16)
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LOCUS 39 (HB3)
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LOCUS 40 (HB5)
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TGTACCTAACCTAAAAAGAAGCCAAGGCAACGAATGTTACCTTGACTTCTAATACATATT
CAACTAATATATATTCAATCATACGCGCATGCGAGAGTGATTGTTGTACATCTATAATG
CGTTGATTTAAAGAACCCTTTATATGGTAAATCAGGTTTGAATAAGTGTTGTATAAATAGA
CCATCTACTAAAACGTCAATGTATGATAATAACTCTCGACGTTCTGTACAATCATTTGCT
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CGAAATGCTTTGACAAGATTTAATGTAATATCCAATTACAAAATGGTTCCGCCACCTAAT
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TCAGTGTATTTCTCGCCATATCTGAACCTTTTGTGAGGCTTTGTTATAACATCCAACACAA
TTAAATGGACATCCTGATACATAAACTGCATCTTACTCCTTCACCGTCAACAAAGCTA
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TAGGCGCTTTCATATGTTTTACTCGTGCGCAATTTCTTTATGACGGCCTTTAATTACTG
GACGTTGAACTGGATTGCCTAGGTAACCACATGTTTCGTTTAACGACATCAACTGTTTTAG
GATTATCATTTGCCACAGTTTCGGGCATTTAAATCCTTTTTTCAGTTGCTTCAAAATCTCCAT
CGTAATCACATTCATAACAATGATC
LOCUS 41 (HB7)
GATCTACATTATATTGCTCAAATAAAGGCGATAATACTTTAGGATTTGGCTTCTCATAGG
CATCCGCTTCGGTAGAAATGATCAAATCGAACAACGAGGTAGCATTGGTATGTGCTAAAA
ATTGTTCTACACCTTTTTTTAGTATCACTCGTAACAATACCAAGTTGATAGCCTTTTGCTT
TCAAATCGATAAGTGCTTCTTTAACACCTTCTACCCAATTAATTTTCAAGGAATACGTTTCA
CTACCAGCTTTTGACTTGTTGACTTGGAACAGTCGGTTGTATCTTGTCCCGTCACATCAT
TAAATGCCTGGATAATTTGTTGTAAAGATCCTGAACCCATCACTGATTTTGGATCAATAG
ATTCTTTAATGACACCGAGTTGTCTTAAAGCAGCTTCTTTATTATGTACTGGGAAAGTCT
CAAGCAATGATTGTACAAATCGTACCCCTATTTTTTCCCAACTTCTATCAAATTCAATTA
ACGTACCATCTTTATCAAATAATATCCATTCCATTGATATCAATACTCCTATTTATTTAT
TTCGTATTATGCTGATTCTATGATATTGTTATCCCCTGAAAATGAACTCGTAGTATTGT
TCTATTTAAATATTGAATTAAATATAATAAAGTGAAATCCCCTTCAATACTTAACAAT
AAACATTGTAACTTAATTTATTACCATGCTTCGCTTCATTGAAAGGGATTTTAGTCATG
ATTAACTTTTGCATATTGTTTTTCATGATTATATTCAATTTTTTATTAATATTTTGGTACAA
CGACTCTCCAACCATTTTTATCTTCTAAAGTACCATTTTGAATACCAGTATAGACGTCGT
ATAATTTTTGAGTAATTTTACCAGTCTCATTATTATTAATAACGATTTTACGATCTTCGT
ATCTCAATGTACCCACAGGTGAAATAACTGCTGCAGTACCACTACCAATACTTCTGTTA
ACTCACCTTTATCATATGATTTCGAATAATTCATCGATTGAAACGCGCGCTCTTCGACTT
CATATCCTAAGTTTTTTAGCTAATTCGATAATAGATTTACGTGTAATACCAGGTAAATAC
TGCCATTCAACTCTGGTGTAATTACTTTGCCATTTTCAACGAAGAAAATGTTTCATGCTAC

CAACTTCTTCGATATATTTCTGTTCAACACCATCAAGCCATAATACTTGGTCATAACCTA
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TAGGACCATAAGGTACTATCTTCAAATCATGCCATCCTTTATCTGCATCATAATCATAAC
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CGTATTAATTTTACAGAGACAAGTAATCTGTGTTTTACTAATATACTTTACATACAAAAA
ACTCTTTACTTTAAAATGAACCTAAGCTCGCGAATTCAATAAGTATAATGAATAATATTAG
AATTCATGCACTAGTTTATTAATAAATAAGAGTAATTTAAAATATCATTCCGTGTATTAAA
GTGAATGGAAATGATTAGTTATTATTTTAAACAGTATCTTTTTGTTCAATAGCTTCTAAC
ATTAATTTAGTCATGCTCGCTAAATCATATTTAGGATC
LOCUS 42 (HB8)
ACGGACTAATATTTCAACTTCCACATTAAAGACACGTTTAATCAACGAATAAAATACGTCT
TGCCGTTGTTGCATTTTCCGTTTGAACATTTATAACAAATTGTTGATTTGAAAGACTAAG
TGCACCATTCATTTCGAATCAGTGCCTGAGCTCTGCTTTTGCATTCATTTTCATCGACGTC
TATTCTAGTTAATTCATTTTTTCATTTCTGATGCAAAGCTCATCGTACAGTCATTCCTTTC
TTATTTAAAACATGATTCACCTTAGAACCCTGTCTATTTTCATTTTTTTTCACAGCTCTA
TTATCATATCATAATATGATTACGTTCTATATTATTTACGTTTATCACTTGGTACGAAAG
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TATGTCTTACTAAATGATTTTTCAGAAATTTCAACTAAATTTGAAGATGTTTTTACATTTA
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CCCCAGGTTGCGTCATCACATTAGAAACATATAGCTTAGGCGCATCAGAATGAATTAACG
CATCTGAAATACCATTTCACACATAAGTTAGAAATAACGCTCGTATATAATGACCCTGGTC
CAAGAACGATTAAATCTGCTTCCCTTAAAGCATCGATTGCTTCTTCCATTGGTTGCACAT
CGTTAGGTTCTAAAACACACGATCAATTTTTTTATGTTTTTTAGGAATATTTGTTTCTC
CAAAAACAATTTCTCCATCTTCCATAACAGCATTTAATTGCACACTTGTATTTGTAGATG
GAATGACTCTACCTTTAATATTTAAAATTTTACTTAATGCTTTAATGGCATGTCCGAAAT
CATTCGTAATATTAGTCATACCTGCGATTAAATAATTACCTAATGAGTGACCGCTAATTT
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TATCATAAAATCTAACAATCAGTTAATTTTTCAAAGAAAATCTCCGTCTCTTTCCATTT
CATAACATAATTATAAACGTCTTTATCTAATCCTGTAAAGGTCTTAAATCTACTACATA
ATATGGATTTGGTAAAATCGTACATCAAATACTAAATCTGCATCCATCTGAATCCCATG
TTTAAAACCGAACTTGTGACATTAATTGTAAAAGTTTCAAACCTTTCATCTTCATAGTA
TCGACGAATGCGTTCTTTAATTCTTTAGGTGATAACTTTGTAGTATCTATAACAAAATT
AGCTATACTTCTAATTTGAGACAAATGCTCTCGCTCATCATTAAATTGCATTGATTAAACGA

TC
LOCUS 43 (HB10)
GATCAACTCATTGCAAAATACGATTTATAGACATCAAAGAATCAATACATTGTAAAGGGG
ATGTTGCCCATGAAAGAAGTTGGATTTGGCACACTAAACTGGGTTGCCGTTATCATTAT
CTACTAGCTATGTTGTTCAATTGGCGTTTATTTTACCAAGCGCGAGCCAAAGTACCAAT
AGTTTCTTTACCGCAAGTGGTCGCTTGCCATCTTGGGTAGTTGGCTTTTCAATTTATGCT
ACTACGTAAAGTGGGATTACATTTATGTCGACACCAGAGAAAGCATTTTTAACAGATTGG
TCATATATCGCTGGTAACATTGCTATCGTCGCAATTATTCCATTACTTATTTATTTCTAT
GTCCCTTTCTTTAAAAAGTTAAAGGTAACATCTGCATATGAATATTTAGAAGCTAGATTT
GGCCCTAGCATACGTGTCATTGGCTCATTATTATTTGTCGTTTACCATTTAGGGCGTGTT
GCAATTGTTATCTACTTACCAACATTAGCAATCACATCTGTATCAGACATGAACCCTTAT
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CCAATTATTTTCTAGGAAATATTTTCAACAACCTTGATCAATACACAGCGAGTCAAGAC
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AATGGTATCCTAGCTTTAATTTACGACCCCTTATTTTATGGTATGGGTACAATGCTGTAT
TCATTTTATACACATGAAGCTGTTTACCAAAAGGCTTCAATACATCATCTGTAGTGCCA
TATTTCATTTTGACTGAGATGCCACCATTGTAGCAGGATTACTTATTGCAGCCATTTTC
GCCGCTGCACAGTCTACCATTTCATCTAGTTTAAATTCTATATCTGCTTGTATTTCAATC
GACATTAAGCAACGCTTCTTCGGAAAAGGTAGCGAGCGACACGAAGTTAACTTTGCACGT
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CCTTTCTATGTATCTACCATTTCATTACAGTTGCTTTTGTCTTTGCTTATATACTTAGC
TTCATTGTCCCTTCAAAACATAAAAAAGATATAACGGGATTAACAATTTTGA AAAAAGAT
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GATTTATAGGGATTGATTACACTGGCGATGATAAGCGCACTAAAAATATTTAACATTACT
GCTGTAACCTACGAACTTGGGTTCAATCATCTGCATATATGAACCTAGCATTGCCATACTA
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GGCAGTAATACGTTTAAAAAGAACGTAAAGCCATTTTATTTTGTATATCTCCAAAAACA
AAATTTATGCCTGCTTTACTA
LOCUS 44 (HD7)

TCCACTCTCTTCGTTGAATCCAAGATTAACGATTGGCAAACAAATTACAGAAGTAATATT
TCAACATAAACGTGTATCTAAATCTGAAGCAAAGTCGATGACAATAGACATTTTAGAAAA
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LOCUS 45 (HD9)
GATCTGAAGTAGCTCGATTTTAAATAGTTTTCAGCAATGACATCGTCTTTTTCTGTCGGC
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ACCTCACCAACACCTTCATGAATTAAAGACATTGGCAATTTCTGAGATAAGACATTCTCA
TCACGGCTACCAGTATAATATCTTTGATC
LOCUS 46 (HE9)
GATCAGATAGATAAAGTATTTTCTTTTTATTATGTTTATCAGAATATGCGCCACCGAAAA
TACCAAATATAATAAATGGAAGTGTGACTCATAACCATCATTGATAATTTTAAAGATG
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TGGTAACTTCGGATCAAATTCGTCTCGATGACCTGGTGTATCGTTTCTGGTCCGTATTC
TGTTAATTCATTAATCGGATC
LOCUS 47 HF6
GATCCAATTGAATTTTTCTCATTACATAAATCTGGATATTGAATGTTAGCAGTTGTT
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TTTAAATACAAAATGTATATCAATATAGTATTACATTTTATAGATAAAGCACAACTTT

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TTGTCTGAGTCCGAATCGCTATCTGAATCTGAGTCGCTGTCTGAATCTGAATCGCTATCC
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CTATCTGAATCTGAGTCGTTGTCTGAGTCCGAATCGCTATCTGAATCTGAGTCGCTATCT
GAGTCTGAGTCGCTATCTGAATCTGAGTCGCTGTCTGAATCTGAATCACTGTCTGAGTCT
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LOCUS 49 (A) B13
TCTTTATTCGAAC TATTAGATTCAC TTTGACCAGTAGTCGTTCCATCAGATCCTTTGTCA
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GGTCCATTAAATTTATATGTAATGTTGTAATGATGGTCATATTTGAATGGCTTTCCATTT
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GAGCAAAC TAAACAGTTCAACCTAAAGTTGAAAAAGTTAAACCTACTGTA ACTACAACA
AGCAAAGTTGAAGACAATCACTCTACTAAAGTTGTAAGTACTGACACAACAAAAGATCAA
LOCUS 49 (B) K16
AGATCAAAC TAAAACACAACTGCTCATACAGTTAAAACAGCACAACTGCTCAAGAACA

AAATAAAGTTCAAACACCTGTTAAAGATGTTGCAACAGCGAAATCTGAAAGCAACAATCA
AGCTGTAAGTGATAATAAATCACAACTAACAAAGTTACAAAACATAACGAAACGCC
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LOCUS 50 (A) GB2
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LOCUS 50 (B) G10

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LOCUS 51 (GC8)
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LOCUS 52 (E1)
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LOCUS 53 (E20)
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LOCUS 54 (E105)
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TTTGTTAATTAAATCGTATAGTATGTCAAGGTTTTGACTTATTGCTTTATTCCGCTCTGG
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LOCUS 55 (E18)
ATCAAAAAGTTATGATGAACGTTTTTACGCCGGATGAAGTAGTCGCATACCAACAACATCA
AGGTAATAAATTTAAAGAACATTTTGATTGGAATTGTTATCTGACACTGCTAGATGTATT
GGATAGTCACAACATTGACCGAGGTCGCACAGACGTAACGCATGTTTTTAAAAATTTAGA
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LOCUS 56 (F5)
AACATACAGGTAAAGTTTTACTTGTAAGTGAAGATAATTTAGAAGGTAGTATTATGTCAG
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LOCUS 57 (F3)
GATCTTCGCGTCTTAATGGATGCCATATACGAACTGAATGACCACCAAGATTTGCGTGAG
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LOCUS 58 (G8)
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AGTCTACTTTTAATGGAACATCTAATTGCAATGCATTTTCCATTATCTCTTCTACAAATT
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LOCUS 59 (G23)
CTTGTAATTCCTTGTTGGTTAAATATGGATGTACCTCAATTTGATTCACCATTGGTTTGA
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LOCUS 60 (G29)
TCTTCTGAGAAGGTTTTTGACCCATTTGCATCATCATCATACGAAGCATTTCCTTCGTTGA
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LOCUS 61A (HA7)
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ACAATTTTGAATCCTACTGGTTTTTGACCTAATTGCTGCAACTGATC
LOCUS 61B (G28)
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CCCACGTGCGATATTTACAAAATCTGCACCTAAACCTAGTGCAATCGCAATTTTATCTGG
TGTCATACTTACCAGATGCCGCCAATTTCACTTTATCTCGAATACCATATTTTCTAA

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LOCUS 62 (H3)
GATCCTTTTGTGTAGACGTAATACGTTCTTGTAATTGTCCCATTTTCAGTAGCAAGTGTT
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LOCUS 63 (GD10)
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CATTTCTATATGCTAATGTGGCAAGATGAGCAAACTCATTTTGTGGATAATGTTTAAAA
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GGAGTTTATCA
LOCUS 64 (F5)
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LOCUS 65 (F110)
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LOCUS 66 (E1)
CAGGATTCGTTTTATCTAACTCTTCCCCAAAAGCTGATAAGTGTGTAGTTTGTGTTG
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LOCUS 67 (F119)
GATCAAAATTTTGAATTAAATACTGTCTCAATTTAAAGTCGAGTTCTTTAAGTGAAATCT
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LOCUS 68 (G27)
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LOCUS 69 (H110)
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LOCUS 70 E100
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LOCUS 71
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LOCUS 72
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LOCUS 73
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LOCUS 74
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LOCUS 75
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LOCUS 76
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LOCUS 77
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LOCUS 78
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LOCUS 83
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LOCUS 85 (F126)
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LOCUS 86
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LOCUS 87
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LOCUS 98 GE2
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LOCUS 99 GE3
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LOCUS 100 GF5
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LOCUS 101 (GF7)
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LOCUS 102 (GF9)
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LOCUS 103 (GF11)
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LOCUS 104 (GF12)
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LOCUS 105 (E18)
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LOCUS 106 (E101)
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LCOUS 107 (E110)
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LOCUS 108 (E125)
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LOCUS 109 (F101)
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LOCUS 113
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NSEVGHMNIAGRIVYQSLTRINKSIEDGDFENDVLNNAIAHVNSHDSALHIFGLLSDG
GVHSHYKHLFALLELAKKQGVEKVYVHAFLDGRDVDQKSALKYIEETEAKFNELGIGQFA
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QNDGVNDGDAVI
LOCUS 15 (I1)
>G2097_STAAU8325, UNDEFINED PRODUCT 1973418:1974263 REVERSE MW:31442
VDLNDRLTFHKRKDRKIVVEIEHNYVP
SNHKNLAYRAAQLFIEQYQLKQGVTSIDKEIPVSAGLAGGSADAAATLRGLNRLFDIGA
SLEELALLGSKIGTDIPFCIYNKTALCTGRGEKIEFLNKPPSAWVILAKPNLGISSPDIF
KLINLDKRYDVHTKMCYEALNRDYQQLCQSLNRLEPISVSKHPQIDKLKNMLKSGAD
GALMSGSGPTVYGLARKESQAKNIYNAVNGCCNEVYLVRLLG
>G2096_STAAU8325, UNDEFINED PRODUCT 1972580:1973401 REVERSE MW:30395
MRYKRSERIVFMTQYLMNHPNKLIPLTFFVKKFKQAKSSISEDVQIIKNTFQKEKLGTVI
TTAGASGGVTYKPMMSKEEATEVVNEVITLLEEKERLLPGGYLFLSDLVGNPSLLNKVGK
LIASIYMEEKLDVVTTIATKGISLANAVANILNLPVVVIRKDNKVTEGSTVSINYVSGS

LOCUS 17 (I3)
>G1894_STAAU8325, UNDEFINED PRODUCT 1776805:1778031 REVERSE MW:45559
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DEIDLKAKDSHGHEYIIDKDAHTRLAKEANTSILRRAFNYVDGTDRTGNFETGLLFIAFQ
KATKQFIDIQNNLGSNDKLNNEYITHRGASASFLVLPGVSKGGYLGETLFD
>G1893_STAAU8325, UNDEFINED PRODUCT 1775112:1776845 REVERSE MW:64202
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LNDALEKAIDNQLSDADAALTHFIETWPYVEGQIQTKDGALYTKIEDKIPYYQSVL
DEHNKAHVKDGLVDLNNQIKEVVGHSSYFVDVMIIFLREGLEVLLIVMTLTMTNRNVKDK
KGTASVIGGAIAGLVLSIILAITFVETLGN SGILRESMEAGLGIVAVILMFIVGVWMH KR
SNKRWNMIKNMYANAISNGNLVLLATIGLISVLREGVEV IIFYMG MIGELATKDFIIG
IALAIVILIIIFALLFRFIVKLIPIFYIFRVLSI
LOCUS 18 (I5)
>G2386_STAAU8325, UNDEFINED PRODUCT 2274220:2275152 REVERSE MW:33616
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PGEQELGGRGVSYCAVCDGAFFKNKRLFVIGGGDSAVEEGTFLTKFADKVTIVHRRDEL R
AQRILQDRAFKNDKIDFIWSHTLKSINEKDGVGSVTLTSTKDGSEETHEADGVFIYIGM
KPLTAPFKDLGITNDVG YIVTKDDMTTSVPGIFAAGDV RDKGLRQIVTATGDGSIAAQSA
AEYIEHLND
>G2387_STAAU8325, UNDEFINED PRODUCT 2275222:2276658 REVERSE MW:57062
HYRLYGIFLLDQLNGKEIVM
TESIWQVLENLNNYEKLYLTYLVQGLTLNKLDFIHRGLLTLYHNELFVSENDVMVAWINQ
GELIIAEKVDLTDVEPYIGAFIYLYFKNQPRNVTKKQITTWLGITQYKLNKMIEFLLSI
LOCUS 19 (I8)
>G2296_STAAU8325, UNDEFINED PRODUCT 2195143:2196150 REVERSE MW:37749
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>G2295_STAAU8325, UNDEFINED PRODUCT 2193368:2195119 REVERSE MW:66415
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FAKEISEKLVVLLPLKFGDYLSSSSMRSLIDIGAPYNHVKVPFAMQSLGALRLTPTRYMK
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AKNDTQQLVSMALAANDRTLYQMICGKDNISKNDVMTLFEDIAQVFLKVTL SFMQYGAL

PELHGQNILLSFEDGRVQKCVLRDHD TVRIYKPWLTAHQLSLPKYVVREDTPNTLINEDL
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>G2294_STAAU8325, UNDEFINED PRODUCT 2192119:2193372 REVERSE MW:44835
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TTMIASPIWGKLGDKISRKWMVLRALLGLAVCLFLMALCTT PLQFVLVRLLOGLFGGVVD
ASSAFASAEAPAEDRGKVLGRLOSSVSAGSLVGPLIGGVTAS ILGFSALLMSIAVITFIV
CIFGALKLIETTHMPKSQTPNINKGIRRSFQCLLCTQQT CRFIIIVGVLANFAMYGMLTAL
SPLASSVNHTAIDDRSVIGFLQSAFWTASILSAPLWGRFND KSYVKS VYIFATIACGCSA
ILOGLATNIEFLMAARILQGLTYSALIQSVMFVVVNACHQ QLKGT FVGT TNSMLVVGQII
GSLSGAAITSYTTPTATTFIVMGVVFVAVSSFLICSTITNQ IND
LOCUS 20 (J7/M10)
>G2187_STAAU8325, UNDEFINED PRODUCT 2068723:2070984 REVERSE MW:85428
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MSVLTITSNVVGKKTGNTPDGRKAGEPFAPGANPMHGRDQ KGALSSLSVAKI PYDCCK
DGISNTFSI VPKSLGKEPEDQNRNL TSM LDGYAMQC GHLNIN VFNRETLIDAMEHP EY
PQLTIRVSGYAVNFIKLTREQQLDVISRTFHESM
>G2186_STAAU8325, UNDEFINED PRODUCT 2067945:2068697 REVERSE MW:28498
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LPYKPYFDASGGGVTVSGGEPLLQMPFLEKLFAELKENG VHTCLDTSAGCANDTKAFQRH
FEELQKHTDLILLDIKHIDNDKHIRLTGKPNTHILNFARK LSDMKQP VWIRHVLVPGYSD
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VNFKGKIPVEL
>G2185_STAAU8325, UNDEFINED PRODUCT 2065846:2067657 REVERSE MW:69718
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LFNMMLKVAGQSQLTINN WTEIVSHPASVILLIIFILSV AF LIYVEFSLLVYMYAGFDR
QIITFKSIFKNAFVNVRKLIGVPVIFVVIYLM LMIPIANLGLSSVLT KNIYIPKFLTEEL
MKTTKGII IYGTFMIAVFILNFKLI FTLP LTILNRQSLFKNMRLSWQITKR NKFR LVIEI
VILELIIGAILTLIISGATYLAICVDEEGDKFLVSSILFV VLKSA LFFYYLFTKLSLISV
LVLHLKQENVLDQPGLEFKYPKPKRKS RFFIISMVLAVT CFIGYNMYLLYNNTINTNISI
IGHRGFEDKGVENSIPSLKAAAKANVEYVELDTIMTKDK QFV VSHDNNLKRLTG VNKNIS
ESNFKDIVGLKMRONGHEAKFVSLDEFIETAKQSNVKLL VELKPHGKEPADYTQRVIDIL
KKHGV EHQYRVM SLDYDVM TKLKKEAPYLKCGYI I PLQFGHFKETSLDFFVIEDFSYSR
LVNQAHLNKEVYTWTINGEEDLT KYLQTNVDGIITDDPALADQIKEEKKDETYFDRSIR
ILFE
>G2184_STAAU8325, UNDEFINED PRODUCT 2065335:2065676 FORWARD MW:12828
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YLNLD AFQKR DILAAHYIAKSAIRTKNLDQMTKAKQRLESIYNSISNPLHSQNN
>G2183_STAAU8325, UNDEFINED PRODUCT 2063238:2065145 REVERSE MW:71718

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NPRTDLKMANFHKYNLEELSMKEYNELQDALKRALDDFHREVKDIKDKNSDLKTFNAEE
DKATKEVYDLVSEIDTLVVSYYGDKDYGEHAKELRAKLDLILGDTDNPHKITNERIKKEM
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VSYGARPTYKKPSKTNAYNVTTHADGTATYGPRVTK
>G2182_STAAU8325, UNDEFINED PRODUCT 2062946:2063050 FORWARD MW:3842
MCVRTRLVSSSSARLSKAIIIAVIVVYHLDVRGLF
>G2181_STAAU8325, UNDEFINED PRODUCT 2061438:2062628 FORWARD MW:42182
MITMQEAYIVAYGRSAAAKAKQGALFHERPDDVAAKVLQGVLRIDGKFNKNMIEDVIVG
TAFPEGLOGQNIARTIALRAGLSDTVPGQTVNRYCSSGLQTIATAANQIMAGQGDILVAG
GVELMSAVPMGGNEPTNPTLQYDDIGASYPMGLTAENVASQFDVSREDQDAYAVRSHQR
AYDAQRDGRFKDEIIPIQVNSVEYTNAGPKVHTNIFDQDEFIRPDTTMEALAKLRTVFKA
DGTMTAGTSAPLSDGAGFVVLMSGDKVKELGVTPIARFVGFKAVGVDPKIMGIGPAYAIP
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TARLLNEMGRRPDSRYGMVTMCIGVGMGAAAI FEYVR
>G2180_STAAU8325, UNDEFINED PRODUCT 2059156:2061414 FORWARD MW:84609
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FDLNLASHLTYGNFDDDLVNDDADLYIEAVKEDIEIKHAVWQQVLQHAKEDALFATNTSG
IPINAIAQAFNEKDQERFFGLHFFNPPRIMKLVELIPTSHTKESIILDVKNFAQNVLGKG
VIVVNDVPGFVANRVGTQTMNDIMYRAEQHKISIVDVALTGQAIGRPKTGTALSDLVG
LDIAVSVIKGMQQVPEETPYFHDVKIVNTLFDNGALGRKTKQGFYKKDKETKARLVYDVE
KQDYVPVSQPQLPILNEFNKDLVHNLDTIFNAQDEAGLFLWETLRNNFYYSAINVPKATD
DFRDIDRALVWGFNWKLGPFQLWDAMGYERVKTRMEDELGDLPOWISDLGDFYKQDETI
EYATPISHFVKDELWDKGDALSVTHDDQLLLKLQSKNNVITDEFNDALVDAIDLLENDH
YTSMVIYADGNFNSVGANLFLMKKAHEDGLVDDVVAQSIDKLHYSFNRLKYS LKPVVTAV
QGRALGGGCELVLYSPIVVAASETYIGLVEAGVGLLPSSGGGLAEMADRILRTSHKFDDKQ
ASMTKVLTNIAFAKVSTNAFEARRYGYLRD TDTIIFNTAQRVEVALKRAKYE AETNYIPN
PRHQYIALGEDFKALIQGQLDAQRRGHFISDHDYHIALNIATILAGGDLPRNTFINQRYI
QSLEKIGFIDLLKSKKSYERIAHMLKTGKPLRN
>G2179_STAAU8325, UNDEFINED PRODUCT 2057714:2058967 FORWARD MW:46482
MHFTLVFILFLGGIYMTFEKETVLKTLFPEDVLSIAKGLTDGEVEFLQQVDSLLESKYRE
NINQHWIDATVPEDYFKDLGELNYFNNPLLYKDRPNAKMPSQLFQFFMSYLLARFDISLA
TLLGVHQGLGHNTFYFGGSKEQIAKYVPKLQSHELRTC FALTEPEHGS DVAGGLETVAER
QGD TWVINGEKKWIGGAHVSDVIPVFAVNKETGKPHCFVVRPEQDGV DIEVIDNKIALRI
VPNALIKLTNVKVDEADRLQNITSFKDIAKILYSTRAGVAYMATGGMAGALRATLDYVTE
RKQFGKPISKYQLIQEKLAMMQGNLAQAMATCAQLANMQAHGEYDEVATSTAKMMNALRL
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LOCUS 21 (G3)
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MNILFAITGIAFALFVAFLE
>G1928_STAAU8325, UNDEFINED PRODUCT 1810990:1811910 REVERSE MW:32866
MANLQKYIEYSREVQQARENNOPIVALESTIISHGMPYPQNVEMATTVEQIIRNNGAIPA
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VIGYQTNELPAFFTRESGVKLTSSVETPERLADIHLTKQQLNLEGGIVVANPIPYEHALS
KAYIEAINEAVVEAENQGIKGDATPFLLGKIVEKTNGKSLAANI KLVENNAALGAKIA
VAVNKLL
G1929
LDHVQQFENASTGSY TALISKEGDMTYGLADMEVFDYITPE
FLIKRSHLLKKAKCIIVDLNLGKEALNFLCAYTTKHQIKLVITTVSSPKMKMNPDSLHAI
DWIITNKDETETYLNLKIESTD DLKIAAKRWNDLGVKNVIVTNGVKELIYRSGEETIIS
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LYHDMEDYKNGKFTKVY
LOCUS 22 (I19)
>G0974 FRG_STAAU8325, UNDEFINED PRODUCT 974673:975977 REVERSE MW:47346
VNEMVNEQIIDISGPLKGEIEVPGDKSMTHRAIMLASLAEGVSTIYKPLLGEDCRRTMDI
FRLLGVEIKEDDEKLVTSPGYQSFNTPHQVLYTGNSGTTTRLLAGLLSGLGIESVLSGD
VSIGKRPM D
>G0975_STAAU8325, UNDEFINED PRODUCT 975981:977042 REVERSE MW:40300
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KVIIPAGEKTKTFEQYQETLEYILSHHVTRNTAIIAVGGGATGDFAGFIAATLLRGVHFI
QVPTTILAHDSVGGKVGINSKQGKNLIGAFYRPTAVIYDLVFLKTL PFEQILSGYAEVY
KHALLNGESATQDIEQHFKDREILQSLNGMDKYIAKGIETKLDIVIADEKEQGV RKFLNL
GHTFGHAVEYYHKI PHGHAVMVGIIYQFIVANALFDSKHDINH YIQYLIQLGYPLDMITD
LDFETLYQYMLSDKKNDKQGVQMVLRQFGDIVVQHVDQLTLQHACEQLKTYFK
>G0976 FRG_STAAU8325, UNDEFINED PRODUCT 977071:978240 REVERSE MW:43249
DFYDSETFKANLDRNDVRVIDDSIAQAMRD KIDEAKNEGDSIGGVVQVVVENMPVGVGSYVH
YDRK
LDGKIAQGVVSINAFKGVSFGEFGKAAEKP GSEIQDEILYNSEIGYYRGSNHLGGLEGGMSN
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LEEFQSNHIEQLKQQIIERRQLNIEF
LOCUS 24:
G0243FRG
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>G0244_STAAU8325, UNDEFINED PRODUCT 218549:220261 FORWARD MW:61780
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RVTRDDVNVADLVISNAVIIDYDKVVKADIGIKNGYIFAIGNAGNPDIMDNVDIIIGSTT
DIIAAEGKIVTAGGIDTHVHFINPEQAEVALESGITTHIGGGTGASEGSKATTVTTPGPWH
IHRMLEAAEGLPINVGFTGKGQATNPALIEQINAGAIGLKVHEDWGATPSALSHALDVA
DEFDVQIALHADTLNEAGFMEDTMAAVKDRVLMYHTEGAGGGHAPDLIKSAAFSNILPS
STNPTLPYTHNTVDEHLDMVMITHHLNAAIPEDIAFADSRIRKETIAAEDVLQDMGVFSM
ISSDSQAMGRVGEVITRTWQVAHRMKEQRGPLDGD FEHNDNNRIKRYIAKYTINPAITHG
ISEYVGSIEPG
>LOCUS 25:
G0027_STAAU8325, UNDEFINED PRODUCT 32103:32513 REVERSE MW:16524
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RHSYELVQGLRDLRLCNWPKFINRLNSVSKKSVSKGVWKVVKYRKHQRMRLNTIYYPA
FNNGAIEGINNKIKLIK
LOCUS 26:
>G2458FRG_STAAU8325, UNDEFINED PRODUCT 2348221:2350185 REVERSE MW:69055
VKIMRVTELLTKDTIAMDLMANDKNGVIDELVNQLDKAGKLSDVASFKEAIHNRESQSTT
GIGEGIAIPHAKVAAVKSPAIAFGKSKAGVDYQSLDMQPAHLFFMIAAPEGGAQTHLDAL
AKLSGILMDENVREKLLHASSPEEV LAI
>G2459_STAAU8325, UNDEFINED PRODUCT 2350185:2351102 REVERSE MW:32573
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SVLDGE
G2460FRG
DRTGCSASTIRRDLSKLQQLGKLQRVHGGAM
LKENRMVEANLTEKLATNLDEKKMIAKIAANQINDNECLFIDAGSSTLELIKYIQAKDII
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LOCUS 27:
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>G1327 STAAU8325, UNDEFINED PRODUCT 1284689:1285450 FORWARD

MW:27870
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NYPGKEDVSVIQVEERAIERGPKGFNFNDNVTPFKYAAGAKAGERIKVIGYPHPYKNKYV
LYESTGPFVMSVEGSSIVYSAHTESGNSGSPVLNSNNELVGIHFASDVKNDDNRNAYGVYF
TPEIKKFIAENIDK
>G1329_STAAU8325, UNDEFINED PRODUCT 1285505:1286227 FORWARD MW:26340
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>G1330_STAAU8325, UNDEFINED PRODUCT 1286327:1287067 FORWARD MW:26652
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VLSVNGNIVTSDAVVQPGSSGSPILNSKREAIGVMYASDKPTGESTRSFAVYFSPEIKKF
IADNLDK
>G1332_STAAU8325, UNDEFINED PRODUCT 1287228:1287941 FORWARD MW:25679
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TGFIVGKNTIVTNKHVVAGMEIGAHIIAHPNGEYNNGGFYKVKKIVRYSQEDIAILHVE
DKAVHPKNRNFKDYTGILKIASEAKENERISVIGYPEPYINKFQMYESTGKVLVSVKGNMI
ITDAFVEPGNSGSAVFNSKYEVVGHVHFGNGPGNKSTKGYGVYFSPEIKKFIADNTDK
>G1333_STAAU8325, UNDEFINED PRODUCT 1288095:1288811 FORWARD MW:25655
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>G1334FRAG._STAAU8325, UNDEFINED PRODUCT 1288994:1290730 FORWARD MW:66904
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SGEDITYQEAWADEEYREDLKAELID
ORF1 (AF7)
SGTGFIVGKNTIVTNKHVVAGMEIGAHIIAHPNGEYNNGGFYKVKKIVRYSQEDIAILH
VEDKAVHPKNRNFKDYTGILKIASEAKENERISVIGYPEPYINKFQMYESTGKVLVSVKGN
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ORF2 (AF7)
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IADNLDK
LOCUS 28 (H130)
>G1388_STAAU8325, UNDEFINED PRODUCT 1337496:1338446 REVERSE MW:36053
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GDFAGNRADSIQVQFKEIKEKMHEKWHEGKYIAYFQAFTNTHAPVEVLKEKFEPVLKEPG
VVGLSIGTRPDCLPDDVVEYLADLNQRTYLWVELGLQTIHQSTSDLINRAHDMKTYDGV
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>G1389_STAAU8325, UNDEFINED PRODUCT 1338556:1339734 FORWARD MW:43345
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LOCUS 41
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>G1926_STAAU8325, UNDEFINED PRODUCT 1808110:1809648 FORWARD
MW:56155
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TTLSAITFMSTPEKAFLTDWSYIAGNIAIVAIIPLLIYFYVPFFKKLKVTSAYEYLEARF
GPSIRVIGSLLFVVYHLGRVAIVIYLPITLATSVDMPNPYIVASLVGLLCILYTFLLGGFE
GVVWSDFIQGVILLGGALVIIILGVVNIKGGFGTVFADAIEHKKLISADNWKLNTAAAI
PIIFLGNIFNNLYQYTASQDVVQRYQASDSLKETNKSLWTNGILALISAPLFYGMGTMLY
SFYTHEAVLPKGFNTSSVVPYFILTEMPPFVAGLLIAAIFAAQSTISSSLNSISACISI
DIKQRFEGKGSERHEVNFAFIIIIAGIFGFGMSLYLIASNSNDLWDLFLFVTGLFGVPL
AGVFAVGIFTKRTNTFGVICGLILGIIFAYVYNGVGKGNSPFYVSTISFTVAFVFAYILS
FIVPSKHKKDITGLTIFEKDKPSTYISKATATK
G1927
>G1927_STAAU8325, UNDEFINED PRODUCT 1809759:1810976 REVERSE
MW:44221
SKAGINFVFGDIQNKNNGFTFFLNVLPLVFIISVLIGIFNYIKVLPFIIKYV
GIAINKITRMGRLESYFAISTAMFGQPEVYLTIKDIIIPRLSRKLYTIATSGMSAVSMAM
LGSYMQMIEPKFVVTAVMLNIFSAIIASVINPYKSDDTDVEIDNLTKSTETKTLNGKTG
KPKKVAFFQMIGDSAMDGFKIAVVVAVMLLAFISLMEAINIMFGSVGLNFKQLIGYVFAP
IAFLMGIPWSEAVPAGSLMATKLITNEFVAMLDKFNVLGDVSARTQGIISVYLVSFANFG
TVGIIVGSIKGISDKQGEKVASFAMRLLLGSTLASIISGSIIGLVL
LOCUS 44

>G2207_STAAU8325, UNDEFINED PRODUCT 2094883:2096472 FORWARD MW:59177
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ESIFKSPQHTYTKRLIDAI PDIHQTRPPRPLNNDILLKFDRVSVDYTSPPSGSLYRAVNDI NLAIRKGETLGIVGESGSGKSTLAKTVVGLKEVSEGFIIWYNELPLSLFKDDELKSLRQEI QMIFQDPFASINPRFKVIDVIKRPLIIHGKVKDNDIIKTVVSLLEKVGLDQTFLYRYPH ELSGGQRQRVSIARALAVEPKVIVCDEAVSALDVSIQKDIIELLKQLQLDFGITYLFIH DMGVINEIC
LOCUS 45
>G2152_STAAU8325, UNDEFINED PRODUCT 2029896:2030945 REVERSE MW:39494
DQRYYTGSRDENVLSQKLPMSLIHEGVGEVVFDSKGVFNKGTKVVMVPNTPTTEKDDVIA
LOCUS 46 G5(1)
>G2647_STAAU8325, UNDEFINED PRODUCT 2528508:2529707 REVERSE MW:44138
VINMLYLEVLKNRNFTYLLIGNFLRRSCFVLFSLQIIWFTVELTNQSSLKLSMMVMSQTL PFIIFGIFGGAYS DKHNKKKILYLS
>G2648_STAAU8325, UNDEFINED PRODUCT 2530085:2534971 REVERSE MW:178787
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LOCUS 47 HF6
>G2560_STAAU8325, UNDEFINED PRODUCT 2436743:2440789 REVERSE MW:146086
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[illegible]

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SMMDTFVKHPIKTGMLNGKKYMMETTNDDYWKDFMVEGQVRVTSKDAKNNTRTIIIFPY
VEGKTLYDAIVKVHVKTIDYDGQYHVRIVDKAFTKANTDKSNKKEQQDNSAKKEATPAT
PSKPTSPVEKESQKQDSQKDDNKQLPSVEKENDASSESGDKDTPATKPTKGEVSSSTT
PTKVSTTQNVAKPTTASSKTTKDVVQTSAGSSEAKDSAPLQKANIKNNDGHTQSQNNK
NTQENKAKSLPQTGEESNKDMTLPLMALLALSSIVAFVLPRKRKN
G1543
>G1543_STAAU8325, UNDEFINED PRODUCT 1497540:1497668 REVERSE MW:4973
MAVPKRRTSKTRKNKRRTHFKISVPGMTECPNCGRIQIITPCM
G1544
>G1544_STAAU8325, UNDEFINED PRODUCT 1497751:1497846 REVERSE MW:3849
MSLLNSKQQDDSESROVDPRLOKLQQLYDKEQ
G1456
>NONE, UNDEFINED PRODUCT 1497815:1498165 REVERSE MW:12767
L...QLVIHITGTYTMPCARLVPVKVPLDVTTEVFDELEGYNQYNDDQDDVDEHYHII
KDGMVNLQDIVEDIVIIEKPMRAYSEQSDQMLTVGNGWEVIDEDQLDELAKQQATR
LOCUS 50 GB2
>G1392_STAAU8325, UNDEFINED PRODUCT 1343118:1349675 FORWARD MW:238192
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YQAQGNVIALGRIHGTDTNDHGDFNGIEKALTVNPNSELIFEFTMTTKNGQGATNVIK
NADTNDTIAEKTVEGGPTLRLFKVPDNRNLKIQFVPKNDAITDARGIYQLKDGKYYSF
VDSIGLHSGSHVFVERRTMDPTATNNKEFTVTTSLKNNGNSGASLDTNDFVYQVQLPEGV
EYVNNSLTKDFPSNNSGVDVNDMNVTYDAANRVITIKSTGGGTANS PARLMPDKILDRLY
KLRVNVNPTPRTVTFNETLTYKTYTQDFINSAAESHTVSTNPYTIDIIMNKDALQAEVDR
RIQQADYTFASLDIFNGLKRRRAQTI LDENRNNVPLNKRVSQAYIDSLTNQMHTLIRSVD
AENAVNKKVDQMEDLVNQNDDELTDDEKQAAIQVIEEHKNEIIGNIGDQTTDDGVTRIKDQ
GIQTLSGDTATPVVKPNAKKAIRDKATKQREIINATPDATEDEIQDALNQLATDETDID
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QQHIAEINANPDATQEERQAAIDKVNAAVTAANTNINANTNADVEQVKTNAIQGIQAIT
PATKVKTDAKNAIDKSAETQHNTIFNNNDATLEEQAQQLLDQAVATAKQNINAADTNQ
EVAQAKDQGTQNIIVVIQPATQVKTDTRNVVNDKAREAITNINATTGATREEKQEAIRVN
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LAQINQHYNALAEINATPDATNDEKNAAINTLNQDRQQAIESIKQANTNAEVDQAATVA
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INQNQTNDQVD
LOCUS 50 G10

>G1392_STAAU8325, UNDEFINED PRODUCT 1343118:1349675 FORWARD MW:238192
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NTNATTEEKQVALNQVDQELATAINNINQADTNAEVDQAQQLGTKAINAIQPNIVKKPAA
LAQINQHYNALAEINATPDATNDEKNAAINTLNQDRQQAIESIKQANTNAEVDQAATVA
ENNID
LOCUS 51 (GC8)
>G2831 FRG_STAAU8325, UNDEFINED PRODUCT 2720353:2721114 FORWARD MW:27865
DPLMLDES�VDIESLSDALMLIESN
>G2832 FRG_STAAU8325, UNDEFINED PRODUCT 2721229:2722446 FORWARD MW:44105
VLRLVEPLKDIDPLNESESLVLVESLIDIESLSEVDSLTLSEPLNDVEVLNEPDVLVEVE
PLVDFESLNESEDSTLSELLSDVDTLNDDESILVLTESLIDCEQLNELDSLTLSDFLNDVE
TLNEPESLTLVEPLIDLESSELDSLTLSESFTDSILCESDMLALITSLADVDVLVESL
NDIDTLIEPDVLALVESDVESLTLSDNDVESLILVDVLVESDILCESLVLVRIEVLVEAD
VLRESLVDVDVLADPDALVLLDVLCESLNDVDVESDSLVLSDVEPDSDVLTVDVKLAMVD
MRFEVDVLSESLNDADVLCESDS
>G2837 FRG_STAAU8325, UNDEFINED PRODUCT 2720004:2726816 REVERSE MW:228019
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DSESQSASAFLESSESTSESTSESVSSSTSESTSLSDSTSESGSTSTSLSNSTSGSTS
ISTSTSISESTSTFKSESVSTSLSMSTSTSLSDSTSLSTSLSDSTSDSKSDSLSTSMSTS
DSISTSKSDSISTSTSLSGSTSESESDSTSSSESKSDSTSMSISMSQSTSGSTSTSTSTS
LSDSTSTSLSLASMNQSGVDSNSASQASNSTSTSTSESDSQSTSSYTSQSTSQSESTS
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TSNNGSASTSTSLSNSASASESDLSTSLSDSTASMQSSESDSQSTASLSDSLSTSTS
NRMSTIASLSTSVSTSESGSTSESTSESDSTSTSLSDSQSTSRSTASGASTSTSTSDS
RSTASTSTSMRTSTSDSQMSLSTSTSTMSDSTSLSDSVSDSTSDSTASTSGMSVS
ISLSDSTSTSTSASEVMSASISDSQMSSESVNDSESVSESNSESDSKMSGSTSVSDSGS
LSVSTSLRKSESVSESSSLSCSQMSDSVSTSDSSLSVSTSLRSSSESVSESDSLSDSKS
TSGSTSTSTSGSLSTSTSLSGSESVSESTSLSDSISMSDSTSTSDSDSLSGSISLSGSTS
LSTSDSLSDSKSLSSSQMSGSESTSTSVSDSQSSSTNSQFDSMSISASESDSMSTSDS
SSISG
LOCUS 52 (E1)
>G0406 FRG_STAAU8325, UNDEFINED PRODUCT 370166:372094 REVERSE MW:70979
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QSTQRMTHEELRVDNODDHSQVSLNGYTKGSEKDQEAFTNNKDEEAVAANKPESEYKVN
EKIKKEHKNFIFGEGVSRGKILAAALLFGMFIAILNQTLNVALPKINTEFNISASTGQWL
MTGFMLVNGILIPITAYLFNKYSYRKLFLVALVLTIGSLICAISMNFPIMMVGRVLQAI
GAGVLMPLGSIVIITIYPPEKRGAMGTMGIAMILAPAIGPTLSGYIVQNYHWNVMFYGM
FIIGIAILIGFVWFKLYQYTTNPKADIPGIIFSTIGFGALLYGFSEAGNKGWGSVEIET
MFAIGIIFIILFVIRELRMKSPMLNLEVLKFPTFTLTIIINMVVMLSLYGGMILLPIYLQ
NLRGFSALDSGLLLPGSLIMGLLGPFAGKLLDTIGLKPLAIFGIAMTYATWELTKLM
DTPYMTIMGIYVLRSGMAFIMPMVTAAINALPGRASHGNAFLNTMRQLAGSIGTAIL

VTVMTTQTTQHLSAFGEELDKTNP
>G0407 FRG_STAAU8325, UNDEFINED PRODUCT 372110:372754 REVERSE MW:23024
MPQKGTIAKLDGMEGSMVQAGNPIAYAYNLLDLYVTANIDEKDIKDVEVGKDVDTIDGQKA SIKGVDSIGKATAASFSLMPSSNSDGNVTKVSQVIPVKITLESEPSKQVVPGMNAEVKIHK N
LOCUS 53 (E20)
>G2244 FRG_STAAU8325, UNDEFINED PRODUCT 2142042:2143301 REVERSE MW:46800
MKLTIVVGLGYIGLPTSIMFAKHGVDVLGVDINQQTIDKLQSGQISIEEPGLQEVYEEVLS SGKLKVSTTPDASDVFIIAVPTPNDDQYRSCDISLVMRALDSILSFLEKGNTIIVESTI APKTMDDFVKPVIENLGFTIGEDIYLVHCPERVLPKGILEELVHNNRIIGGVTEACIEAG KRVYRTFVQGEMIETDARTAEMSKLMENTYRDVNIALANELTKICNNLNINVLDVIEMAN KHPRVNIHQPGPGVGGHCLAVDPYFIIAKDPENAKLIQTGREINNSMPAYVVDTTKQIIK VLSGNKVTVFGLTYKGDVDDIRESFAFDIYELLNQEPDIEV
>G2245_STAAU8325, UNDEFINED PRODUCT 2143358:2144242 REVERSE MW:33683
MRKNILITGVHGYIGNALKDKLIEQGHQVDQINVRNQLWKSTSFKDYDVLIHTAALVHNN SPQARLSDYMQVNMLLTKQLAQKAKAEDVKQFI FMSTMAVYGKEGHVGKSDQVDTQTPMN PTTNYGISKKFAEQALQELISDSFKVAIVRPPMIYGAHCPGNFQRLMQLSKRLPIIPNIN NQRSALYIKHLTAFIDQLISLEVTGVYHPQDSFYFDTSSVMYIEIRRQSHRKTVLINMPM LNKYFNKLSVFRKLEGNLIYSNTLYENNNALEIIPGKMSLVIADIMDETTTKDKA
>G2246_STAAU8325, UNDEFINED PRODUCT 2144245:2144799 REVERSE MW:21063
MKRLFDVVSIIYGLVVLSPILLITALLIKMESPGPAIFKQKRPTINNELFNIYKFRSMKI DTPNVATDLMDSTSYITKTGKVIKTSIDELPQLLNVLKGEMSIIVGPRPALYNQYELIEK RTKANVHTIRPGVTGLAQVMGRDDITDDQKVAYDHYLTHQSMMLDMYIIYKTIKNIVTS EGVHH
>G2247 FRG_STAAU8325, UNDEFINED PRODUCT 2144813:2146015 REVERSE MW:46577
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LOCUS 54 (E105)
>G2254 FRG_STAAU8325, UNDEFINED PRODUCT 2152390:2153505 REVERSE MW:42140
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>G2255_STAAU8325, UNDEFINED PRODUCT 2153408:2155321 REVERSE MW:72361
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FGKNVDIVPIIADVQNRARMFEIMETYKPYAVYHAAAHKHVPLMEDNP EEAVRNNILGTK
NTAEAAKNAEVKKFVMISTDKAVNPPNVMGASKRIAEMI IQSLNDETHRTNFVAVRFGNV
LGSRGSVIPLFKSQIEEGGPVTVTHPEMTRYFMTIPEASRLVLQAGALAE GGEV FVLDMG
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>G2256_STAAU8325, UNDEFINED PRODUCT 2155251:2156012 REVERSE MW:29362
DQLFFELQSKGFVPPIIAHPERNKAI SQNL DILYDLINKGALSQVTTASLAGISGKKIRKLAI
QMIENNLTHFIGSDAHNTEIRPFLMKDLFNDKKLRDYYEDMNGFISNAKLVVDDKKIPKR
MPQQDYKQKRWFGL
LOCUS 55 (E18)
>G2912 FRG_STAAU8325, UNDEFINED PRODUCT 2797518:2798504 FORWARD MW:37832
SKSYDERFTPDEVVAYQQHQGNKFKEHFDLNCYLTL LDVLD SHNIDRGRTDVTHVFKNLETK
VLTMGFIDDLLYPDD
LOCUS 56 (F5)
>G1261 FRG_STAAU8325, UNDEFINED PRODUCT 1216923:1217903 FORWARD MW:36061
HTGKVLLVTEDNLEGSIMSEVSAIIAEHCLFDLDAPIMRLAAPDVPSM
PFSPVLENEIMMNPEKILNKMRELA EF
>G1262_STAAU8325, UNDEFINED PRODUCT 1217919:1219190 FORWARD MW:46726
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NGRFSPVVFKLASEHDIDLSQVVGSGFEGRVTKDIMSVIENGTTAQSDKQVQTKSTSV
DTSSNQSSDNSENSTIPVNGVRKAIAQNMVNSVTEIPHAWMMIEVDATNLVNTRNHYKN
SFKNKEGYNLTFFAFFVKAVADALKAYPLLNSSWQNEIVLHKDINISIAVADENKLYVP
VIKHADEKSIKGIAREINTLATKARNKQLTAEDMQGGTFTVNNTGTFGSVSSMGIINHPQ
AAILQVESIVKKPVVINDMIAIRNMVNLCISIDHRILDGLQTGKFMNHIKQRIEQYTLEN
TNIY
>G1263_STAAU8325, UNDEFINED PRODUCT 1219532:1219978 FORWARD MW:16676
VIELMDMNF DLYMNGVVEQARNEIESAGYEQLTTAEDVDKVLKQDGTTLVMINSVCGCAG
GIARPAASHALHYDVLPDRLVTVFAGQDKEATQRAREYFEGYAPSSPSFALVKDGKITEM
IERHQIEGHDVMNVINQLQTLFNKYCEER
>G1264_STAAU8325, UNDEFINED PRODUCT 1219995:1220972 FORWARD MW:36973
MLKLNPKYKIGFRTIKTAVGMTLGVIISKLLGLDNYASSAILVVLCIKHTKVHSLQAIISR
LVSCFLVLFLGSAIFSLGQSPIVLGIIVLLFIPLTVVLKVQEGVITSCVILLHVFN AKS
IDAHLIVNETLLLLIGLSIAFTMNLMMPSLDKQLDEYKCKIEQQIADIFSKYSYICEKYE
DTIAIEFEVLLLNKKAKSIAFRDVKNHFVRNENSYYHYFDMREEQVELLMRMKPLIESI
CHKD
LOCUS 57 (F3)

>G0451_STAAU8325, UNDEFINED PRODUCT 410768:412549 FORWARD MW:67976 DLRVLMDAIYELNDHQDLREITKDSKMOKLALAGFLKKIKGTYIESLLKEHKLL
>G0452_STAAU8325, UNDEFINED PRODUCT 412872:414536 FORWARD MW:60909 MEMSVTEVIFSFLGGLGIFLYGLKIMGDGLQASAGDRLRDILNKFTSNPVLGVIAGIVVT ILIQSSSGTTVITIGLVTAGFMTLKQAIGVIMGANIGTTVTAFIIGIDLGEYAMPILALG AFLIFFFKRSKINNIGRILFGFGSLFFGLEFMGDAVKPLASLDGFKQLMLDMSTNPILAV IVGAGLTALVQSSSATIGILQEFYQODLISLNAAPVLLGDNIGTTITAILASLAGSIAA KRAALVHVIFNLIGVIIFTIFLPVVIHLISLLQDLWHLKPAMTIAVSHGIFNITNTLIQL PFVAGLAWIVTKLVPGKDIADDYKPOHL
LOCUS 58 (G8)
>G0922_FRG_STAAU8325, UNDEFINED PRODUCT 915062:915931 REVERSE MW:33411 MPPELPEVEHVKRGIEPYVINQKIEHVIFSDKVIEGKAQGKETIIKGIELDTFKTLSEGYT ITNVERRSKYIVFQLDNKREQRTLISHLMAGGFFIVDELEDIMIPNYRKHHVIFELSN DKKLIYSDIRRFGEIRNVASVASYPSEFLEIAPEPFSNEALTYLNRHQQSNKNKPIKQV IL
>G0923_FRG_STAAU8325, UNDEFINED PRODUCT 915950:918577 REVERSE MW:99163 DELIFEVPKSEVDSFSEFVEEIMENALQLDVPLKVDSSYGATWYDAK
LOCUS 59 (G23)
>G2454_FRG_STAAU8325, UNDEFINED PRODUCT 2344101:2344937 REVERSE MW:32360 MLNEIQILNNGYPMPVGLGVYKISDEDMTKVVNAIDAGYRAFDYFYDNEASLGRAL KDNQVDREDLFITTKLWNDYQGYEKTFEYFNKSIENLQTDYLDLFLIHWPCADGLFLET YKAMEELYEQGKVKAIGVCNFNVHHLEKLMAQSSIKPMVNQIEVHPYFNQQLQ
>G2455_STAAU8325, UNDEFINED PRODUCT 2345162:2346508 REVERSE MW:51133 LETSTIISLIIFILLIALTTVFVGSEFALVKIRATRIEQLADEGNKPAKIVKKMIANLDY YLSACQLGITVTSGLGLWLGEPTFEKLLHPIFEAINLPTALTTTISFAVSFIIVTYLHV LGELAPKSIAIQHTEKLALVYARPLFYFGNIMKPLIWLNMNGSARVIIRMFGVNPDAQTD MSEEEIKIIINNSYNGGEINQTELAYMQNIFSFDERHAKDIMVPTQMITLNEPFNVDEL LETIKEHQFTRYPIITDDGDKDHIKGFINVKEFLTEYASGKTIKIANIYIHELPMISETTRI SDALIRMQREHVHMSLIIDEYGGTAGILTMEDILEEIVGEIRDEFDDDEVNDIVKIDNKT FQVNGRVLLDDLTEEFGEFDDSEDIDTIGGWLQSRNTNLQKDDYVDTTYDRWVSEIDN HQIIVILNYEFNEARPTIGQSDEDEKSE
LOCUS 60 (G29)
>G0139_FRG_STAAU8325, UNDEFINED PRODUCT 137065:137352 REVERSE MW:11080 VMNLAKFSRIKKAGETMATWVAIIFIVAALILGLIGGFLARKYMMDYLLKKNPPINEEML RMMMMQMGMQKPSQK

>NONE, UNDEFINED PRODUCT 137582:139645 REVERSE MW:75349
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SKTPGHPEYRHTDGVVTTGPLGQGFAMSVGLALAEDHLAGKFNKEGYNVVDHYTYVLAS
DGDLMEGISHEAASFAGHNKLSKLVVLYDSNDISLDGELNKA FSENTKARFEAYGWNYYL
VKDGNDLEEIDKAITTAKSQEGPTII EVKTTIGFGSPNKAGTNGVHGAPLGEVERKLTFE
NYGLDPEKRFNVSEEVYEIFQNTMLKRANEDESQWNSLLEKYAETYP ELAEFFKLAI SGK
LPKNYKDELPRFELGHNGASRADSGTVIQAI SKTVPSFFGGSADLAGSNKSNVNDATDYS
SETPEGKNVWFGVREFAMGA AVNGMAAHGGLHPYGATFFVFS DYLPALRLSSIMGLNAT
FIFTHDSIAVGEDGPTHEPIEQLAGLRAIPNMNVIRPADGNETRVAWEVALESESTPTSL
VLTRQNLPLVDVPEDVVEEGVRKGAYTVYGSEETPEFLLL ASGSEVSLAVEAAKDLEKQG
KSVRVVSM PNWNAFEQQSEYKESVIPSSVTKRVAIEMASPLGWHKYVG TAGKVIAIDGF
GASAPGDLVVEKYGFTKENILNQVMSL
LOCUS 61 (G28/HA7)
>G2610_FRG STAAU8325, UNDEFINED PRODUCT 2494989:2495441 FORWARD MW:17293
DLGMDKDEAKKLF AKSESIFKDLKGVKYKVDYKDKKAIEHLDIDYTEVDMKKLNKRLGV
STKENKDISFEKLEKQLKHRGLKEKDKMDDK
>G2611_STAAU8325, UNDEFINED PRODUCT 2495615:2497207 REVERSE MW:58937
LGGGIVMTFLTVMQFIVNIIVVGFM LTVIVIGLIWLIKDKRQS QHSVLRNYPLLARIYI
SEKMGPELRQYLFSGDNEGKPF SRNDYKNIVLAGKYNSRMTSFGTTKDYQDGFYIQNTMF
PMQRNEISVDNTTLLSTFIYKIANERLFSREEYRVPTKIDPY YLSDDHAIKLGEHLKHPF
ILKRIVGQSGMSYGALGKNAITALS KGLAKAGTWMNTGEGGLSEYHLKGNGDIIFQIGPG
LFGVRDKEGNFSEGLFKEVAQLSNVRAFELKLAQGA KTRGGHMEA EKVNEEIAKIRNVEP
YKTINSPNRYEFIHNAEDLIRFVDQLQQLGQKPVGFKIVVSKVSEIETLVRTMV ELDKYP
SFITIDGGEGGTGATFQELQDGVGLPLFTALPIVSGM LEKYGIRDKVKLAASGKLVTPDK
IAIALGLGADFN IARGMMISVGCIMSQQCHMNTCPVGVATTD AKKEKALIVGEKQYRVT
NYVTSLHEGLFNIAAAVGVSSPTEITADHIVYRKVDGELQTIH DYKCLKLIS
LOCUS 62 (H3)
>G2004_STAAU8325, UNDEFINED PRODUCT 1871545:1872954 REVERSE MW:51401
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AFDELSTEVQILETG IKVVDLLAPYIKGGKIGLFGGAGVGKTVLIQELINNIAQEHGGIS
VFAGVGERTREGNDLYFEMSDSGVIKKTAMVFGQMNEPPGARMRVALSGLTMAEYFRDEQ
GQDVLLFIDNIFRFTQAGSEVSALLGRMPSAVG YQPTLAT EMGQLQERITSTTKG
LOCUS 63 (GD10)
>G2900_FRG STAAU8325, UNDEFINED PRODUCT 2781950:2783308 FORWARD MW:51966
DPIFKQEVERNLEKEIRNV
>G2901_STAAU8325, UNDEFINED PRODUCT 2783589:2784719 FORWARD MW:41914

MMEFTIKRDYFITQLNDTLKAISPRITLPILTGIKIDAKEHEVILTGSDSEISIEITIPK
TVDGEDIVNISETGSVVLPGRFFVDI IKKLPGKDVKLSTNEQFQTLITSGHSEFNLSGLD
PDQYPLLPQVSRDDAIQLSVKVLKNVIAQTNFAVSTSETRPVL TG VNWLIQENELICTAT
DSHRLAVRKLQLEDVSENKNVIIPGKALAE LNKIMSDNEEDIDIFFASNQVLFKVG NVNF
ISRLLLEGHYPDTRLPENYEIKLSIDNGEFY
LOCUS 64 (F5)
>G1261 FRG_STAAU8325, UNDEFINED PRODUCT 1216923:1217903 FORWARD MW:36061
HTGKVLVLTEDNLEGSIMSEVSAIIAEHCLFDLDAPIMRLAAPDVPSM
PFSPVLENEIMMNPEKILNKMRELAEF
>G1262 STAAU8325, UNDEFINED PRODUCT 1217919:1219190 FORWARD MW:46726
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AGQTVAIDTIICKIETADEKTNETTEIQAQVDEHTQKSTKKASATVEQTSTAKQNQPRN
NGRFSPVVFKLASEHDIDLSQVVGSGFEGRVTKKDIMS VIENG GTTAQSDKQVQTKSTSV
DTSSNQSSDNSENSTIPVNGVRKAIAQNMVNSVTEIPHAWMMIEVDATNLVNTRNHYKN
SFKNKEGYNLTFFAFFVKAVADALKAYPLLNSSWQGNEIVLHKDINISIAVADENKLYVP
VIKHADEKSIKGIAREINTLATKARNKQLTAEDMQGGTFTVNNTGTFGSVSSMGIINHPQ
AAILQVESIVKKPVVINDMIAIRNMVNLCISIDHRILDGLQTGKFMNHIKQRIEQYTLEN
TNIY
>G1263 STAAU8325, UNDEFINED PRODUCT 1219532:1219978 FORWARD MW:16676
VIELMDMNF DLYMNGVVEQARNEIESAGYEQLTTAEDVDKVLKQDGTTLVMINSVCGCAG
GIARPAASHALHYDVLDPDRLVTVFAGQDKEATQRAREYFEGYAPSSPSFALVKDGKITEM
IERHQIEGHVDMNVINQLQTLFNKYCEER
>G1264 STAAU8325, UNDEFINED PRODUCT 1219995:1220972 FORWARD MW:36973
MLKLNPKYKIGFRTIKTAVGMTLGVIISKLLGLDNYASSAILVVLCKIKHTKVHSLQAIISR
LVSCFLVFLGSAIFSLGQSPIVLGIIIVLLFIPLTVVLKVQEGVITSCVILLHVFN AKS
IDAHLIVNETLLLLIGLSIAFTMNLMMPSLDKQLDEYKCKIEQQIADIFSKYSYICEKYE
DTIAIEFEVLLLNIKKAKSIAFRDVKNHFVRNENSYYHYFDMREEQVELLMRMKPLIESI
CHKD
LOCUS 65 (F110)
>G2848 STAAU8325, UNDEFINED PRODUCT 2734525:2735082 REVERSE MW:21969
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NIIMMNQNSNYSIDALYQFLFEFIFDIEERYIRMYVQLSNTPEEFSGNIYGQIQDLNQS
LSKEIAKFYDESKIKMTKEDFQNLILLFLESWYLKASF SQKFGAVEESKSQFKDEVYSLL
NIFLKK
>G2849 STAAU8325, UNDEFINED PRODUCT 2735246:2736481 FORWARD MW:47752
LQFFNLLFYFVFMSTIYWIWGSYFYFTREIRYSLNKKPDINVDELEGITFLLACYNESE
TIEDTL SNVLALKYEKKEIIIINDGSSDNTAELIYKIKENND FIFVDLQENRGKANALNQ
GIKQASYDYVMCLDADTIVDQDAPYYMIENFKHDPKLGAVTGNPRIRNKSSILGKIQTIE
YASLIGCIKRSQTLGAVNTISGVFTL FKKS AVVDVGYWDTDMITEDIAVSWKLHLRGR

IKYEPLAMCWMLVPETLGGLWKQVRWAQGGHEVLLRDFSTMKTKRFPLYILMFEQIIS
ILWVYIVLLYLGYLFITANFLDYTFMTYSFSIFLLSSFTMTFINVIQFTVALFIDSRYEK
KNMAGLIFVSWYPTVYWIINAAVVLVAFPKALKRKKGGYATWSSPDRGNTQR
>G2850_STAAU8325, UNDEFINED PRODUCT 2736448:2736750 FORWARD MW:11783
MVKPRQREYPTLKSSLNIVRETALIAISCVFWIYCLVLLVYIGTIFEIHDESINTIRVA
LNIENTEILDIFETMGIFAIIFVFFTISILIQKWQRGRES
>G2851_STAAU8325, UNDEFINED PRODUCT 2736729:2737619 FORWARD MW:34958
MAERKRIVKYRKFIILVLSILILPVSTLDGHHIANADDDSPKKLKYKENSALALNYHRV
RKANFLNNFIYFFSSSKEIKNYSVSQSQFESQIKWLKSHDAKFLTLKEFLYKKGKFKPK
RSVWINFDDMDETIYENAYPILKKYKIPATGFIITGHVGEENFHNLDMISKKELKEMYKT
GLWEFETHTHDLHNLSKNNKSKLMKASEATIIKDLNKSEKYLTKNFKKSQKTIAYPYGLM
NDDKLPVIKKAGLKYGFSLEEKAVTPNSNDYIIPRILISDDAFEHLIKRWDGFHEKD
>G2852_STAAU8325, UNDEFINED PRODUCT 2737609:2738658 FORWARD MW:41344
MKKIRLELVYLRAIICAIIIIITHLLTQITLKHENMEGGSLVLQFYIRNIVIFGTPCFIIL
SQLLTTLNYQKVTRYRLTTRVKYILIPYILMGLFYSYSESLLTDSSFNKQFIENVLLGQW
YGYFIVVIMQFFILSYIIFKINYNLFNSKILLLSFILQQSFLYYFTNNTAFHDTVLHYY
PLSENTIIFGWIFYFFLGAYMGYNYERVLNFLERYLVIMIVLAVATYFVFIALANGDYWN
VTSFSYSLTPYNSIMFIVILGICTHFKTMLFNTIQMISAFSFFIYLLHPIILDSLFAITN
IFEDNTMVFLAISLLFILGLCIGVMILREFYIFRFIIGKQPYKLNINAY
>G2853_FRG STAAU8325, UNDEFINED PRODUCT 2739111:2741162 REVERSE MW:77120
DPIVLVHGFNGFTDDINPSVLAHYWGGNKMNIHQDLEENGYKAYEASISAFGSNYD
RAVELYYYIKGGRVDYGAHAHAKYGHERYGKTYEGIYKDWKPGQKVHLVGHSMGGQTIHQ
LEELLRNGNREEIEYQKKHGGGEISPLFKGNHDNMISSITTLGTPHNGTHASDLAGNEALV
RQIVFDIGKMFNGKNSRVDFGLAQWGLKQKPNESYIDYVKRVKQSNLWKSNDNGFYDLTR
EGATDLNRKTSLNPNIVYKTYTGEATHKALNSDRQKADLNMFFPFVITGNLIGKATEKEW
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LOCUS 66 (E1)
>G0406_STAAU8325, UNDEFINED PRODUCT 370166:372094 REVERSE MW:70979
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EKIKKEHKNFIFGEGVSRGKILAAALLFGMFIAILNQTLNVALPKINTEFNISASTGQWL
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GAGVLMPLGSIVIIITIYPPEKRGAMGTMGIAMILAPAIGPTLSGYIVQNYHWNVMFYGM
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MFAIGIIFIILFVIRELRMKSPMLNLEVLKFPTFTLTIIINMVVMLSLYGGMILLPIYLQ
NLRGFSALDSGLLLPGSLIMGLLPGFAGKLLDTIGLKPLAIFGIAVMTYATWELTKLNM
DTPYMTIMGIYVLRSGMAFIMPMVTAAINALPGRLASHGNAFLNTRQLAGSIGTAIL
VTVMTTQTTQHLSAFGEELDKTNP
>G0407_STAAU8325, UNDEFINED PRODUCT 372110:372754 REVERSE MW:23024
MPQKGTIAKLDCMEGSMVQAGNPIAYAYNL

DDLYVTANIDEKDIKDVEVGKDVDVTIDGQKASIKGKVDSIGKATAASFSLMPSSNSDGN
YTKVSQVIPVKITLESEPSKQVVPGMNAEVKIHKN
LOCUS 67 (F119)
>G1831 FRG_STAAU8325, UNDEFINED PRODUCT 1723090:1723806 REVERSE MW:27770
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KELKNVTGYRYSKGGKHLYLFDKNRKFTRVQIFGKDIERFKARKNPGLDIFVVKEAENRN
GTVFSYGGVTKKNQDAYDYINAPRFQIKRDEGDGIATYGRVHYIYKEEISLKELDFKLR
QYLIQNF
>G1832_STAAU8325, UNDEFINED PRODUCT 1724158:1725096 REVERSE MW:34671
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KPNATTPPSTKVEAPQQTANATTPPSTKVTPPSTNTPQPMQSTKSDTPQSPTTKQVPTE
INPKFKDLRAYYTKPSLEFKNEIGIILKKWTTIREFMNVVPDYFIYKIALVKGDDKKYGE
VHRNVDVFFVLEENNYNLEKYSVGGITKSNSKKVDHKAGVRITKEDNKGTISHDVSEFKI
TKEQISLKELDFKLRKQLIEKNNLYGNVSGSKIVIKMKNNGGKYTFELHKKLQENRMADVI
DGTNIDNIEVNIK
>G1834_STAAU8325, UNDEFINED PRODUCT 1725193:1725327 REVERSE MW:5264
LFVKVAFCLCLKSDETSNVPSVESHQNHFYLTNIMDFLIYLTMIQI
>G1835_STAAU8325, UNDEFINED PRODUCT 1725449:1726531 REVERSE MW:40775
LEHTIMKMRTIAKTSALGLLTGAVTTTQSVKAEKIQSTKVDKVPTLKAERLAMINIT
AGANSATTQAANTRQERTPKLEKAPNTNEEKTSASKIEKISQPKQEEQKTLNISATPAPK
QEQSQTTESTTPKTKVTPPSTNTPQPMQSTKSDTPQSPTIKQAQTDMPKYEDLRAYY
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DNKYQLKKYSVGGITKTNSKKVNHKVELSITKKDNQGMISRDVSEYMITKEEISLKELDF
KLRKQLIEKHNLYGNMGS GTIVIKMKNNGGKYTFELHKKLQEHMADVIDGTNIDNIEVNI
K
>G1837_STAAU8325, UNDEFINED PRODUCT 1726810:1727562 REVERSE MW:28926
DYDFFPFKIDKEAMSLKEIDFKLRKYLIDNYGLYGEMSTGKITVKKKYYGKYTFELDKKLQE
DRMSDVINVTD
IDRIEIKVIKA
LOCUS 68 (G27)
>G0516_STAAU8325, UNDEFINED PRODUCT 482272:486597 REVERSE MW:163057
VVIVLAMTEQQKFKVLADQIKISNQLDAEILNSGELTRIDVSNKNRTWEFHITLPQFLAH
EDYLLFINAIEQEFKDIANVTCRFTVTNGTNQDEHAIKYFGHCIDQTALSPKVKGQLKQK
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EAHIQEEDEQSARLATEKLEKMKAEKAKQQDNNE SAVDKCQIGKPIQIENIKPIESIIEE
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VRAQGRIEEDTFIRDLVMMMSDIEEIKKATKKDKAEKRVFHLHTAMSQMDGIPNIGAY
VKQAADWGHPAIAVTDHNVVQAFPDAAHAAAEKHGKMIYGMELVDDGVPIAYKPQDVV
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TDDMLVDAPEIEEVLTEFKEWVGDAIFVAHNASFDMGFIDTGYERLGFGPSTNGVIDTLE
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NKKLSNEDAYKRARPSHVTLIVQNQQGLKNLFKIVSASLVKYFYRTPRIPRSLLDEYREG
LLVGTACDEGELFTAVMQKDQSQVEKIAKYDFIEIQPPALYQDLIDRELIRDTETLHEI
YQRLIHAGDTAGIPVIATGNAHYLFEHDIARKILIASQPGNPLNRSTLPEAHFRTTDEM
LNEFHFLGEEKAHEIVVKNNTNELAD
LOCUS 69 (H110)
>G2217 FRG_STAAU8325, UNDEFINED PRODUCT 2108154:2110211 FORWARD MW:74420
DPASGYASILGIPTLQTGVFGGIIIGALAAWCYNKFYNINLPSYLGFFAGKRFPVIMM
ATTSFILAFPMALIWPTIQSGLNAFSTGLLDSNTGVAVFLFGFIKRLLI PFGLHHIFHAP
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AKPENKKVVAGLMGSAALTSFLTGTGITEPLEFSFLFVAPLLFFIHAVLDGLSFLTLYLLDL
HLGYTFSGGFIDYFLLGILPNKTQWWLVI PVGLVYAVIYYFVFRFLIVKLKYKTPGREDK
QSQAATASATELPYAVLEAMGGKANIKHLDACITRLRVEVNDKSKVDVPGLKDLGASGVL
EVGNMQAIFGPKSDQIKHEMQQIMNGQVVENPTTMEDDKDET VVVAEDKSATSELSHIV
HAPLTGEVTPLSEVPDQVFSEKMMGDGIAIKPSQGEVRAPFNGKVQMIFPTKHAIGLVSD
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LOCUS 70
>G1778_STAAU8325, UNDEFINED PRODUCT 1669401:1669715 REVERSE MW:11597
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AVDPDDIEMLODLVLAATNEAMNKADEL TQERLGKHTQG
>G1780_STAAU8325, UNDEFINED PRODUCT 1669808:1671502 REVERSE MW:63481
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AINCLNSTDGEPCNECHICKGITQGTNSDVIEIDAASNNGVDEIRNIRDKVKYAPSESKY
KVYIIDEVHMLTTGAFNALLKTLEPPAHAIFILATTEPHKIPPTIISRAQRFDFKAISL
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DTEYRALMNLELDMLYQMIDLINDTLV SIRFSVNQNVHFEVLLVKLAEQIKGQPQVIANV
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QIAKVLDKANKADI KLLKDHQWEVIDHAKNNKKSLSVSLQNSEPVAA SEDHVLVKFEEE
IHCEIVNKDDEKRSSIESVVCNIVNKNVKVGVPSDQWQVRTEYLQNRKNEGDDMPKQQ
AQQT DIAQKAKDLFGEETVHVIDEE
>G1781_STAAU8325, UNDEFINED PRODUCT 1671574:1672095 REVERSE MW:19908
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DVVGHVLLIEVEINSDDKTYYG LAIASLSVHPELRGQKLGRGLVQAVEERAKAQEYSTVV
VDHCFDYFEKLG YQNAAEHDIKLESGDAPLLVKYLWDNLTDAPHGIVKFPEHFY
>G1782_STAAU8325, UNDEFINED PRODUCT 1672236:1672334 REVERSE MW:3948
LKTIQRIIRGTCLWEVAFLYVKFDSSEL DVQFE

>G1783_STAAU8325, UNDEFINED PRODUCT 1672737:1673480 REVERSE MW:28585
IGNDVASDSIYDYLEKVLNL
NISYSSKSITFEPFDEQAYQLFGDVSVAYSATVRSIVYLENTMPFQYNISKHLANEFKEN
DFSRRRIK
LOCUS 71
>G1083_STAAU8325, UNDEFINED PRODUCT 1057165:1058778 REVERSE MW:57664
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LOCUS 72
>G2296_STAAU8325, UNDEFINED PRODUCT 2195143:2196150 REVERSE MW:37749
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IIAMPSHIGGEHAISGIKWIGSKHDNPSKRNMERASGVIIILNDPETNYPIAVMEASLISS
MRTAAVS VIAAKHLAKKGFKDLTIIGCGLIGDKQLQSMLEQFDHIERVVFVYDQFSEACAR
FVDRWQQORPEINFIATENAKEAVSNGEVVITCTVTDQPYIEYDWLQKGAFI
>G2297_STAAU8325, UNDEFINED PRODUCT 2196150:2197127 REVERSE MW:35879
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EPDAHGGYLMTRIAKVQELLATIDDAYWINQYANELNWQSHYHGAGTEIVETIKQPIDYF
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SEINQVIHVDDYQSALGCRKLIDYEGIFAGGSTGSI IAAIEQLITSIEEGATIVTILPDR
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LOCUS 73
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SLDMGTVVAGTKYRGFEERLKKVMEEIQQAGNVILFIDELHTLVGAGGAEG AIDASNIL
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AGWTGIPLTKINETESEKLLSLEDTLHERVIGQKDAVNSISKAVRRARAGLKD PKRPIGS
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QLTEKVR RKPYSVILFDEIEKAHPDVFNILLQVLDDGHLTDTKGRTVDFRNTIIIMTSNV
GAQELQD
LOCUS 74
>G1438_STAAU8325, UNDEFINED PRODUCT 1399373:1401364 REVERSE MW:74364
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NSSQLSHQNI VSMIDVDEEDDCYLLVMEYIEGPTLSEYIESHGPLSVD TAINFTNQILDG
IKHAHDMRIVHRDIKPQNILIDSNKTLKIFDFGIAKALSETSLTQTNHVLGTVQYFSPEQ

AKGEATDECTDIYSIGIVLYEMLVGEPPFNGETAVSIAIKHIQDSVPNVTTDVRKDIPQS
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TPNTGERVERGDSVDVVISKGPEKVKMPNVIGLPKEEALQKLKSLGLKDVITIEKVYNNQA
PKGFIANQSVTANTEIAIHDSNIKLYESLGKQVYVEDFEHKSFSKAKKALEEKGFKVES
KEEYSDDIDEGDVISQSPKGKSVDEGSTISFVVSKGKSDSSDVKTTTESVDVPYTGKND
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>G1439_STAAU8325, UNDEFINED PRODUCT 1401364:1402104 REVERSE
MW:28046
DQLMQLALDNHSDNVTFILA
AIEGDKV
LOCUS 75
>G0364_STAAU8325, UNDEFINED PRODUCT 331693:334395 REVERSE
MW:98970
MAANFKEQSKKHFDLNGQSYTYDLKAVEEQGITKVSNLPSYIRVLLLESLLRQEDDFVIT
DDHIKALSQFGKDGNEGEVPFKPSRVILODFTGVPVAVDLASLRKAMDDVGGDITKINPE
VPVDLVIDHSVQVDSYANPEALERNMKLEFERNYERYQFLNWATKAFDNYNAVPPATGIV
HQVNLEYLASVVHVRDVGDKTAFPDTLVGTDSHTTMINGIGVLGWGVGGIEAEAGMLGQ
PSYFPIPEVIGVRLVNSLPQGATATDLALRVTLQELRKKGVVGKFEFFGPGVQHLPLADR
ATIANMAPEYGATCGFFPVDDDESLKYMKLTGRSDEHIALVKEYLKQNHMFFDVEKEDPNY
TDVIELDLSTVEASLSGPKRPQDLIFLSDMKSSFENSVTAPAGNQGHGLDKSEFDKKA EI
NFKDGSKATMKTGDIATAAITSCTNTSNPYVMLGAGLVAKKAVEKGLKVPEYVKTS LAPG
SKVVTGYLRDAGLQPYLDDLGFNLVGYGCTTCIGNSG
LOCUS 76
>G2434_STAAU8325, UNDEFINED PRODUCT 2324870:2325844 REVERSE
MW:37506
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>G2435_STAAU8325, UNDEFINED PRODUCT 2326069:2327847 REVERSE
MW:68170
HGLMKGYTTSELSHLIDELRFKGFLENDEI
LMCDTSIKKLLSNEVEVFTTPFKQKATEKVFINTVEGVDRVLFSQ LVEVRKKLSDKLTIA
PVSIFSDYTLEEFKRKPASKQDMINIDGVGSYKLKHYCPAFLETIQNYKAKV
LOCUS 77
>G2617_STAAU8325, UNDEFINED PRODUCT 2501985:2502917 REVERSE
MW:34781
DRAIRSVAFFLTALPSYWIASILIIYVSVKLNILPTSGLTGP

LOCUS 78
IIAIIILIFISFFFSGSETALTAANKAKFKTEADKGDKKAKGIVKLLLEKPSEFITILIG NNVANILLPTLVTIMALRWGISVGIASAVLTVVIILISEVIPKSVAAATFPDKITRLVYPI INICVIVFRPITLLLNLKLTDSINRSLSKGQPQEHQFSKEEFKTMALAIAGHEGALNEIETS RLEGVINFENLKVKDVDTPRINVTAFASNATYEEVYETVMNKPYPYTRYPVYEGDIDNIIG VFHSKYLLAWSNKKENQITNYSAPLFFVNEHNKAEWVLRKMTISRKHLAIVLDEFGGTEA IVSHEDLIEELLGMEIEDEMDKKEKEKLSQQQIQFQQRKNRNVSI
LOCUS 79
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MEROZOITE SURFACE ANTIGEN
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SURFACE PROTEIN
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LOCUS 81
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LOCUS 82
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LOCUS 103 (GF11)
>G2235 FRG_STAAU8325, UNDEFINED PRODUCT 2133494:2134471 REVERSE MW:36941
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>G2236 STAAU8325, UNDEFINED PRODUCT 2134482:2135219 REVERSE MW:28095
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LOCUS 104 (GF12)
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>G2829 FRG_STAAU8325, UNDEFINED PRODUCT 2717099:2718649 REVERSE MW:61259
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LOCUS 105 (E18)
>G2912 FRG STAAU8325, UNDEFINED PRODUCT 2797518:2798504

FORWARD MW:37832
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>G1083 FRG_STAAU8325, UNDEFINED PRODUCT 1057165:1058778 REVERSE MW:57664
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LOCUS 107 (E110)
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LOCUS 108 (E125)
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LOCUS 109 (F101)
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>G1100_STAAU8325, UNDEFINED PRODUCT 1071126:1072409 REVERSE MW:46849
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>G1101_STAAU8325, UNDEFINED PRODUCT 1072584:1072829 REVERSE MW:9040
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LOCUS 110 (F113)
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LOCUS 111
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G2821
>G2821_STAAU8325, UNDEFINED PRODUCT 2706470:2707033 REVERSE MW:20989 SDDKHDFIIEQILSRSCDIESVESWKSSL
LOCUS 112
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LOCUS 113
G1111
>G1111_STAAU8325, UNDEFINED PRODUCT 1083909:1085690 FORWARD MW:65093 DPSEINKVIHVDLGIIADCKRFLECLNDKNVETIEHSDWVKHCQNNKQKHPFKLGEEDQVFC KPQQTIEYIGKITNGEAIVTTDVGQHQMWAQFYPPKHNHGQWVTSGGLGTMGFGIPSSIGAK LANPDKTVVCFVGDGGFQ MTNQEMALLPEYGLDVKIVLINNGTLGMVKQWQDKFFNQRFSSHVFNGQPDFMKMAEAYG VKGFLIDKPEQLEEQLDAAAFAYQGPALIEVRISPTEAVTPMVPSGKSNHEMEGL
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>G1112_STAAU8325, UNDEFINED PRODUCT 1085693:1085944 FORWARD MW:9621 MTRILKLQVADQVSTLNRITSFAFVRLQYNIDTLHVTHSEQPGISNMEIQVDIQDDTSLHI LIKCLKQQINVLTVECYDLVDNEA
G1113
>G1113_STAAU8325, UNDEFINED PRODUCT 1086069:1087085 FORWARD MW:37588 LEEFIMTT
LOCUS 114
G1542
>G1542 STAAU8325, UNDEFINED PRODUCT 1495403:1497337 FORWARD

MW:72192
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G1543
>G1543_STAAU8325, UNDEFINED PRODUCT 1497540:1497668 REVERSE MW:4973 MAVPKRRTSKTRKNKRRTHFKISVPGMTECPNCGRIQIITPCM
G1544
>G1544_STAAU8325, UNDEFINED PRODUCT 1497751:1497846 REVERSE MW:3849 MSLLNSKQODDSESRQVDPRLQKLQQLYDKEQ
G1546
>NONE, UNDEFINED PRODUCT 1497815:1498165 REVERSE MW:12767 DQDDVDEHYHIIKDGMVNLQDIVEDIVIIEKPMRAYSEQSDQMLTVGNGWEVIDEDQLDELA KQQATR
LOCUS 115
G2712
>NONE, UNDEFINED PRODUCT 2598712:2601288 REVERSE MW:94980 EVGDRYYNRTIIYTVYLNYNVDFKRRQYTLAKFLYKMGTFIAKHKWSAVIAWIVIVAAIL IPLATNAPKFDNDIKMTGLESOLDTNKKIEKHFNQDSEKAQIRVVFKTTKDDGIVQPNITK DIKKTLDIDIKDDKHIDKISD
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>G2713_STAAU8325, UNDEFINED PRODUCT 2601346:2601891 FORWARD MW:21879 MKETDLRVIKTKKALSSSLQLLEQQLFQTITVNQICDNALVHRTTFYKHFYDKYDLLEY LFNQLTKDYFARDISDRLNHPFQTMSTINNKEDLREIAEFQEEAEFNKVLKNVCIKIM HNDIKNNRDRIDIDSDIPDNLFYIYDSLIEGFIHWIKDEKIDWPGEDIDNIFHRLINIK IK
G2714
>G2714_STAAU8325, UNDEFINED PRODUCT 2601974:2602138 REVERSE MW:6456 VRYVISIIMGIVLAIWSFKQLSQSHLD SGFIFFIVYVLCISCFNSDKHDKNKKR

G2715

>G2715_STAAU8325, UNDEFINED PRODUCT 2602253:2603800 REVERSE
MW:57130

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TABLE 9 DNA SEQUENCES STAPHYLOCCOCUS EPIDERMIDIS

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LOCUS 4:
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LOCUS 5:
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LOCUS 7:
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LOCUS 8 :
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LOCUS 9:

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LOCUS 10:
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LOCUS 11:
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TCGAATTCTATTCTATTCTTTTCTATACAATGAAACGGGTGTCATACATCATCAGTAACT
AATTATGATAGATATGAACTTGTGGTCTTTATCGTCTTTAGTTTTACTAATGAGAGCAC
GTGGAGTATTTCCATCTTTGATTCTAATTTCACTCATCTAGTTTATCAAAATATTTTT
CGGCTTGCTCTGTAACATATTGTGTAATACCTATCGTTTCTGCCTGTCCGTAATAATCTA
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CTTGTGTGAAGTTTGAAAAATACGATTGCAAATTATCATTAACTGTTTAAAGTTATTAT
TCAGCGTTTCATCATAATCAGCTGCAGTTGACGAAGGAATTAAGGCTGCTTTTTTCATTAA
TATTATCCCAAGAGTTAATTTTAGTTTTACCCTCTTCAACCGTAGTACCAACTATAAATT
CACCTGGTGTAAATGGAATCTTGACTTGATTGTTTATAGATAGCAAATGAATAGGAATAT
CTTTCAAATCACTATTCTCACGTAAACGAGAAAGCATTTCACTAGCCATCTGTTTACCTT
GCTTTTCAATCTCTTTATCAGATAAATCTTTACTAAATGTTTCGCCATCTTTCTCTTTTT
TGTAATAATAAACACTATTTCATGGCTAAACCAATTGTCATCCCTTTTATATTTTTTACCTT
TAGAATCACTATTTCCATAAAAATCCTGCTCGAGTATATTTGAAAGATAGGCTGGAGAAT
TTTCAGCTATTTTCTCTTCATCTGTTTCACCATTGTGAGATGGATTGAGTCCAAGATTCT
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AATACTTATTTGTTGGAAGATTTCTTTACTTAATTCTAGTAAACCACTTTCAAATCTT
CTCCATTATAACCATTGTCATATTATCTTGTAATAATCCACGAGCCTGGCTTTCTTTGA
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GCTTCATTACGTACTCCTCTAATTATTAGATTCCATTTTGTTTTTCAATAAATGCTGCT
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TTCAACCATTGAGATGCTTCATTTCTCGTCATTTGCTCGAGTTTCCCTGTTAATACAATT
GTTTTCCCACTAAAATCAGGATGACCTTCGATTTGAGTTGTTTAAATTCCTTTATAAGAC
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TCAGTTACTTTAAAAAGTTGATC
LOCUS 12:
GATCCTGACACAGCTATTTCTCTCTTAGATAATC
CTATTCAACCTTTACCTAATAATAAGAAAGTATAATTAGATACATCAAAGGGGCAATCT
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TCTGCATTAGTCGGTGCTCTTTTTATTGCTAATTACTTTGGTAATATTCAATTATCTAAA
TACGAATTGTTACAACCTAGCGACTGAAATTGAGGGACACCCTGATAATGTAGCACCTACA
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AGAATAGAAGTTCCGCACGTAGATATAATTTTAACTATACCTCCATATGAGCTTCGTACA
GAAGACTCTAGAAGGGTCTTACCCGATACATTTTCACATAAAGGTGCTGTGCAAAATAGT
GCCATTAGTAACACTATGATTTGTGCTCTCATTGAGCATAAATATAAACTTGCTGGAAAG
ATGATGGAACAAGATGGTTTTTCATGAACCATATAGGCAACACCTTATTCAGAATTCAAT
CAAGTACGTAACTATCACGTCAACATGATGCATATGCAACTGTTATCAGTGGAGCTGGA
CCTACAATACTCACTCTTTGTCCAAAAGAAAAAGTGGTAAATTAGTTAGAACACTACGT
GAGAAAATTAATAATTGTGCTTCAGAACTAGTAACAATTAATGAAATAGGTGTTAAAGAT
GAAGTGGTGTACCTAAAGTCCTAAATTATTGTAAAATATAGTTAAGAATAAACTTTTAAAT
AACTCTTGAAAGGAGTTCTATACTATATGACTCAGTATAAAATGGTAGTTTTAGATATGG
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ATATTCAAAGCGTGGTTATTATGTAGTATTGGCCTCAGGTAGACCAACAGAAGGTATGT
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GAGGTAAACTATAAATATGGCTAATGAAAATGTAGAGGTGATCAGCCTGTTTCAAAGG
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 CCGGATTACCTATGAATCGTGTGCTGATTGGAAGGAATATATTAATCATAGTGTGCCCA
 AAGTTATGGGTGTGGATTATGTAGGTCATATTACCGAAGCACGTATTGAATTGGATGGTT
 ACTTCAATAATGATATTGATGTGACAACGAGTAAGCCTTTTTTCTAGAGTTTATGGCAA
 AGAATGTTTCGAAGGGGAACGCAATAAAAGCACTTTGTAAAAGATTACAAATTTCTCTAG
 AAGAAGTTATAGTATTCGGGGACAGTTTGAATGATAAGTCAATGTTTGAAGTTGCTGGAT
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 TAGATAATAATTCTAACGGTATTCCTTATGCTTTAAAAGAACTTTTGCTTTAAAGTATTA
 TTACAATGAATTAATATGTAAATTAATAATTTTAAGGTTAATTGAATCTGACTTCTCTAA
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 TACAATTTCTTTATTGAATATATTTAAATCTTTATTTACTTTTCTTTCAAATCAATTGA
 AAATCGAGACTTTCAATTGATTTGCTATTTTCGAGTATGTGTCCAGTCATGTTTTCCTTT
 ATAGCGTTTAACATGTGCATATACTTGATC

TABLE 10 PROTEIN SEQUENCE STAPHYLOCOCCUS EPIDERMIDIS

LOCUS 1:

ORF1:

DQTALKQAEKAKSEVTQSTTNVSGTQTYQDPTQVQPKQDTQSTTYDASLDEMSTYNEISS
 NQKQOSLSTDDANQNQTNVTKNQEEETNDLTQEDKTSTDNTNLQETQSVAKENEKDLGA
 NANNEQQDKMTASQPSENQAIETQTASNDNESQOKSQQVTSEQNETATPKVSNTNASGY
 NFDYDDEDDSSDHLPIISLNNVNATSKQTTSYKYKEPAQRVTNTNVKKETASNQATID
 TKQFTPFSAQAQPRTVYSVSSQKTSSLPKYTPKVNSSINNYIRKKNMKAPRIEEDYTSYF
 PKYGYRNGVGRPEGIVVHDTANDNSTIDGEIAFMKRNYTNAFVHAFVDGNRIIETAPTDY
 LSWGAGPYGNQRFINVEIVHHDYDSFARSMNNYADYAATQLQYYNLKPDSEAENDGRGT
 WTHAAISNFLGGTDHADPHQYLRSHNYSYAELYDLIYEKYLIKTKQVAPWGTTSTKPSQP
 SKPSGGTNNKLTVSANRGVAQIKPTNNGLYTTVYDSKGHKTDQVQKTLSTVTKTATLGNNK
 FYLVEDYNSGKKYGWVKQGDVVYNTAKAPVKVNQTYNVKAGSTLYTPWGTTPKQVASKVS
 GTGNQTFKATKQQQIDKATYLYGTVNGKSGWISKYLLTASKPSNPTKPSTNNQLTVTNN
 SGVAQINAKNSGLYTTVYDTKGKTTNQIQTLSVTKAATLGDKKFYLVGDYNTGTNYGWV
 KQDEVIYNTAKSPVKINQTYNVKPGVKLHTVPWGTYNQVAGTVSGKGDR

LOCUS 2:

ORF1:

RIGGKYMDNIKIIVASDSIGETAELVARAGVSQFNPQCKHEFLRYPYIESFENVDEVIQ
 VAKDTNAIIVYTLIKPEIKKYMISKVNEHALKSVDIMGPLMELLSNSIEETPYEPPGMVH
 RLDDAYFKKIDAIEFAVKYDDGKDR

ORF2:

GEAFMVKNMDTIVQLAKHRGFVFPGSDIYGGLSNTWDYGPLGVELKNNIKKAWWQKFITQ
 SPYNVGIDAAILMNPKTWEASGHLGNFNR

ORF3:

RPIELSQRQEQUIEIVKSEGPITGEHIAEKINLTRATLRPDLAILTMSGFIEARPRVGYF
 YSGKSKNKIINEKLRKYVVKDYMSHPVVIKENMTVYDAICTIFLEDVSTLFITNENNDFV
 GVCSRKDLLRASMIGEDIHTMPISVNMTRMPHVSYLKEQELVIYAANQMIDKEIDSLPIV
 RPKENDKFEVIGRISKTTITKLFVSLFKE

LOCUS 3:

ORF1:

SVMKNFILSVQHLLAMYAGAILVPIIVGTSLKFSAEIAYLVTVDIFMCGVATFLQANKV
TGTGLPIVLGCTFTAVAPMILIGQTKGLDVLYGSLISGILVVLIAFFFSYLVKFFPPVV
TGSVVTIIGINLMPVAMNYLAGGEGAKNYGDTKNLILGGVTLLIILILQRFKGLKSIA
ILIGLAIGTALAGIFGMVDIKQVGDAHWFGFPVPRFSGFGFDVSSILVFFIVAVVSLIE
STGVYHALSEITGRKLERKDFRKGytaEGLAILGSIFNAFPYTAYSQNVGLVSLSGAKK
NNVIYGMVILLIICGCI PKLGALANI IPLPVLGGAMIAMFGMV MAYGV SILGNINFQONQ
NLLIIAISVGLGAGISAVPQAFKGLGEQFAWLTQNGIVLGAISAIILNFFFNGIKYKQTE
ENVK
ORF2:
VESLGRKVKEDGVVIDEKILKVDGFLNHQIDAKLMNDVGKTFYESFKDAGITKILTIEAS
GIAPAIMASFHFDVPCLFAKKAKPSTLKDGfySTDIHSFTKNKTSTVIVSEEFLGADDKV
LIIDDFLANGDASLGLNDIVKQANATTVGVGIVVEKSFQNGRQRLEDAGLYVSSLCKVAS
LKGNKVTLLGEA
ORF3:
NWRLFLMWENKFAKESLTFDDVLLIPAASDVLPSDVL SVKLSDKI
LOCUS 4:
ORF1:
YWTYHFKEKGMVIMDDLKQONQSSNEKPKGNKIINILIFIGMILLIQIPIGVSLIALPFS
VKFSKLTSLIALSMLITGTALLI IWLVRNYLSHTYERQYQSMRGKDIFINIGFLVLSMVF
SILSSVLMVIFTGNDTTANEKEINESL DLLLQKDHLPHISIVATVVL MICIIGPYLEELL
FRGIFKETLFMKYRFWLPFIISII FSSQHLSTNIFSYAIYFLMGCVLYLAYNRRRNKID
SMMVHMLNNSVSTLPV FVGYLWLYFR
ORF2:
DLHI IKGDTPEVKSHTTLGHEGIGIEEIGDNVNNFKVGDKVIISCISSCGKCYCKGI
YAHCEGGGWILGHLVNGTQAEYVKVPFADNSLYHAPSNLKEDALVMSDILPTGYEIGV
LKGKVKPGCTVAIVGAGPVGLAALLTAQFYSPSKIIMIDLDDNRLETAKELGATHLINSK
ETETAIKKVKSLNPRGVDVAIEAVGIPQTFDLQNLIGVDGTIANVG VHGLPVQLDIDKL
WIKNINVTTLGLVSGNTTEELLEALKSKI IQPEQLVTHYSKLSEIESAYDLFRNATDHKAI
KLIENDITI
LOCUS 5:
ORF1:
QIVQRKGCHLMKIRVIVPCYNEGEVVLKTYDKLTEIMKKDSL IKNYEYDLLFINDGSTDT
TIHHIKNIVAYDNHV KYLSFSRNFGKEAAMIAGYQHSTMHDAVIMIDGDLQHPPEYIPQM
IEGYIEGYDQVVAKRNRQGENFVRKTL SRCYKLI NAFVEDIQFEDGVGDFRLLSRRAVQ
ALTTLDEYNRFSKGLFEWIGYETKVFOYENV TREDGESKWTFRKLLNYGIDGLISFNKPK
LRMMIYLG MFTFSISILYIIYLLINILINGINIPGYFTTIAAILLLGGIQLMSIGVVGEY
IGRIYYEVKHRPKYIVENSNIQTENLDMRYNALNLNKNRNNKRSNDLYKLSSFYKVKTYS
DTYASNYSQDEGFKERVH
ORF2:
DQLLVN ILQPYEQHIKQENRTLEVNFCTDIDAFYQYRPPIERILTNLLDNALKFSNSGSR
IDIIISECKENDVISISIKDEGIGIVPELQSRIFERTFRVEDSRNTKTGGSGLGLYIANE
LAQQIDASITVQSDLDIGTTMTLTLKKFQFKK
LOCUS 6:
ORF1:
SIAGAAIASQGSFAVLHYQGFTKIIIVLII SPIIAFCVGYMMYTIVKIVFKNSNLRTNR
NFRFFQIFTAALQSFSHGTNDAQSMGIITLALIVGNLQDGSNVEPQVWVKVACATAMGL
GTAVGGWKIIKTVGGNIMKIRPANGAAADISSALTIFVASSLHFPLSTTHVSSSILGVG
ASNRAKGVKWSAQRMVVTWVITLPISAVLA AIIYFI IHLFLK
ORF2:
GGVTLKKLAFAITAASGAAVLSHHDAAEASTQHKVQSGESLWTIAQQYNTSVESIKQNNN

LSNNMVFPQVINVGGSASQNTSSNTSSSSASSHTVVAGESLNI IANKYGVSDALMQAN
HLNGYLIMPNQILTIPNGGSGSGSGGTATQTSNGYTSPSFNHQNLYTEGQCTWYVFDKRS
QAGKPISTYWSDAKYWASNAANDGYQVDNTPSVGAIMQSTPGPYGHVAYVERINGDGSIL
ISEMNYANGPYNMNYRTIPASEVSSYAFIH
LOCUS 7:
ORF1:
DHIIRAYHKFLQSGYQTELHLFGRDEDNQIPLMNTLISELKLSDKVKIFKYTNQPLQEFK
NSKASLLTSQYEGFGLTLMESIEMGCPVLSYNVRYGPSEIIQNGINGYLIEKNDIDSLSK
HMINIIIEHPLQKVKNKDTLKYNAAVNNYKQLMQSLDLLK
ORF2:
SRGGFQVQKKYITAIIGTTALSALASTHAQAATHTTVKSGESVWSISHKYGISIAKLKSL
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NNYLIFPGQKLKVSGKATSSSRKASGSSGRTATYTVKYGDSLAIASKYGTTYQKIMQL
NGLTNFFIYPGQKLKVPGGSSSSSSSNNTSRNGGYSPTFNHQNLTYWGQCTWHVFNRRRA
EIGKGISTYWWNANNWDNASAADGYTIDYRPTVGSIAQTDAGYYGHVAFVERVNSDGSIL
VSEMNWSAAPGNMTYRTIPAYQVRNYKFIH
LOCUS 8:
ORF1:
DQFREAMTKFPVWMGATTLFFGAINGAKEMLDVITEIDGKMITLAKVTGDDNALQQTFFID
ANNAASQFGQTLGSLVDVYAEFARQGVKGNELSQFSNAALIAANVGEIDAKQASEYLTSM
SAQWETTGNQAMRQVDSLNEVSNNKYATTVEKLAQQQAKAGSTAKSMGLTFDETNGIIGAL
TAKTKQSGDEIGNFMKATLPKLYSGKGKSTIEGLGISMKDENGQLKSAISLLEEVSQKTK
NLEKDQKAAVINGLGGTYHYQRMQVLLDDLSKTDGLYKQIKESSESSAGSALQENAKYME
SIEAKVNQAKTAFEQFALAVGETFAKSGMLDGIRMVTQLLTGLTHGITELGTTAPIFGMV
GGAASLMSKNVRSGFEGARSSVANYITEVNKLAKVNNAAGQVVGLQKVQGTASQLQFNK
NGEYDKAASQAKAAEQATYQFSKAQKDVSAAMIASGAINKTTVATTASTVATRAATLAV
NGLKLAFRGLLAATGVGLAITGVSEFVLEKVVGSFNAASQAAEQYKQKQEQTKQAIASMSN
GEINSLISSYDKLQKMNNSGSAFNTEAEAEKYKEVTSQLANIFPDLVTGENRYGKEMAGNK
EVMKQKIELIKQEMELERQKNAIKQKEEQDAYIKEQDSLAKKNRGQKWYQLGQTPCLKLO
EQARPTTVSDNSNINKINATIQKVKSQAQAEKALEQVDKQLAQSQTKNRQNEVQHLQKVR
QALQDYITKTGOANQATRAAVLTAQQQFTNQIATMKKLGTGQQVMTTISNSVAKTAKSG
KAAQATFKSFETSLVKSSSFKSKMASYEASVKKFKNAANQSAKIAALKDVERDYSKVAKG
IMQAAKAANMSKSKMKDLKKSLOQNIQAETGFRASVSKAGKVTIDQSKKIKQNR
LOCUS 9:
ORF1:
VLWGVFDMDLLIGTLFLILVLVIFTLFTYKAPSGMRAMGALANAALIASFLVEAFNKYVGG
QVFGIKFLEELGDAAGGLGGVAAAGLTALAIGVSPVYALVIGAACGGMDLLPGFFAGYIV
GYMMKYTEKYVPDGIDLIGSIILLAPIARLIATGLTPVNNTLIKIGDIIQSSTDANPLI
MGIVLGGIITVVGTAPLSSMALTALLGLTGAPMAIGAMAAFFSSAFMNSALFHLRLKLGDRK
STISVGIEPLSQADIVSANPIPIYVTNFFGGAIAAGIIIAWSGMINNATGTATPIAGFLVM
FGFNSLTKVIIYGVVMAIIGTIAGIVGSIIVFKKYPITKKQMLERDTT
LOCUS 10:
ORF1:
MEIKQIKYFVEVVRQGGMTQASEHLYIAQSTISKAIKNIENEYDITLFDRSQKQIKLTDI
GQTFYDNSLEFLALFEKLSLEMNDIVNVQKGIKIGLSPMMNVQMFTNALNQFHRLYPNV
TYEVIEGGGKIVENLTSNDDVDIGITTLPVDL
ORF2:
LSESANSFYLVHDDFLIRIVKECLLTHVNSKLMMLWRFVMSGFFNRMTRKENPTIYQNKDG
HLKRTLVRDFLALGVGTIVSTSIFTLPGVVAAEHAGPAVALSFLLAIVAGLVAFITYAE
MASTMPFAGSAYSWINVLFGEFGWVAGWALLAEYFIAVAFVASGFSANLRGLIAPLGIS

LPKSLSNPFGSNGGVIDIIAAVVIILTALLSRGMNEAARMENVLVILKVLAIILFVIVG
LTAINFSNYIPFIPEHKVTETGDFGGWQGIYAGVSMIFLAYIGFDSIAANSAEAINPQKT
MPRGILGSLIVAIVLFVAVALVLVGMFHYSQYADNAEPVGWALRESGHGIIAAIVQAISV
IGMFTALIGMMLAGSRLLYSFGRDGLLPSWLSQLNKHKLPNRALVILTIIGVVIGSR
LOCUS 11:
ORF1:
DPETLFIVMSQILFHLPLVGGFLLAAILAAIMSTISSQLLVTSSSLTEDFYKLIRGSDKAS
SHQKEFVLIGRLSVLLVAIVAITIAWHPNDTILNLVGNWAGFGAAFSPLVLYSLYWKDL
TRAGAIISGMVAGAVVIVWISWIKPLATINAFFGMYEII PGFIVSVLITYIVSKLTKKPD
DYVIENLNKVKHVKE
ORF2:
DQLFKVTESEELIEIQDIGDKLAQSVVTYLENSDIRSLIEKLSNKNVNMSYKGIKTTEIEG
HPDFSGKTIVLTGKLEQMTRNEASEWLKMQGAKVTNSVTKSTDIVIAGADAGSKLAKAEK
YGTEIWTEAAFIKQNGI
ORF3:
MKRTIFLLMSILLLLTACGDGHKQTSSDKEQSEHKDNHNKNQVKQIATDKKVQGDNYRTI
LPPKESQARGLLQDNMANGYNGEDFESGLLELSKEIFPTNKYLYQDGQYLDKKTINAYLD
PKYTKKEIDKMSEKEKKSKNANENLGLNPSHNGETDEEKIAENSPAYLSNILEQDFYGNS
DSKGKNIKGMTIGLAMNSVYYYKKEKDGETFSKDLSDKEIEKQKGQMASEMLSRLRENSD
LKDIPHFAYKQSSQDSITPGEFIVGTTVEEGKTKINSWDNINEKAALIPSSAADYDE
TLNNNFQFNDNLQSYFSNFTQAVGKVKFVNKKAKQLTVDLPIDYYGQAETIGITQYVTE
QAEKYFDKLDEYEIRIKDGNTPRALISKTKDDKEPOVHIYHN
LOCUS 12:
ORF1:
LDTSGQSSMEEVLKLKIPASTANLGVGFDSIGMALDKYLHMSIRKIERANWEFLYYSSE
LEGLPKDENNYIYQTALNVARKYNVTLPQLQIEMRSDIPLARGLGSSASALVGALFIANY
FGNIQLSKYELLQLATEIEGHPDNVAPTIYGGIAGFYNPITKITDVARIEVPHVDIILT
IPPYELRTEDSRRVLPDTFSHKGAVQNSAISNTMICALIQHKYKLAGKMMEQDGFHEPYR
QHLLIPEFNQVRKLSRQHDAYATVISGAGPTILTLCPKEKSGKLVRTLREKINNCASELVT
INEIGVKDEVVYLS
ORF2:
LLKGVLYYMTQYKVVLDMDDTLMNSDNKLSIETKSYLLDIQKRGYYVVLASGRPTEGML
PTARELELNKYNSFII SYNGGKTINMANENVEVDQPVSKEDFDNIVDYCRDKNFLVLTID
NGYIIHDSHEYMNIESQLTGLPMNRVADLKEYINHSPKVMGVVDYVGHITAEARIELDGY
FNNDIDVTTSKPFLEFMAKNVSKGNAIKALCKRLQISLEEVIVFGDSLNDKSMFEVAGY
SVAMGNASDELKKIADDEVTLDNNSNGIPYALKELLV

CLAIMS

1. An antigenic polypeptide, or part thereof, encoded by an isolated DNA
5 molecule selected from the group consisting of:
 - (i) DNA molecules represented by the DNA sequences in Table 7 or 9;
 - (ii) DNA molecules which hybridize to the sequences identified in (i) which
encode a polypeptide expressed by a pathogenic organism; and
 - (iii) DNA molecules which are degenerate as a result of the genetic code to the
10 DNA sequences defined in (i) and (ii),
for use as a vaccine.
2. An antigenic polypeptide according to Claim 1 wherein said DNA molecule
is genomic DNA.
15
3. An antigenic polypeptide according to Claim 1 or 2 wherein said DNA
molecule hybridizes to the the sequences in Tables 7 or 9 under stringent
hybridization conditions.
- 20 4. An antigenic polypeptide according to any of Claims 1-3 wherein said
polypeptide (s) are represented by the amino acid sequences in Tables 8 or 10.
5. An antigenic polypeptide according to any of Claims 1-4 wherein said
polypeptide is derived from a bacterial genus/species selected from the group
25 consisting of: *Staphylococcus spp.*; *Staphylococcus aureus*; *Staphylococcus*
epidermidis; *Enterococcus faecalis*; *Mycobacterium tuberculosis*; *Streptococcus*
group B; *Streptococcus pneumoniae*; *Helicobacter pylori*; *Neisseria gonorrhea*;
Streptococcus group A; *Borrelia burgdorferi*; *Coccidioides immitis*; *Histoplasma*
sapsulatum; *Neisseria meningitidis type B*; *Shigella flexneri*; *Escherichia coli*;
30 *Haemophilus influenzae*.

6. An antigenic polypeptide according to Claim 5 wherein said polypeptide is derived from the genus *Staphylococcus* spp.
7. An antigenic polypeptide according to Claim 6 wherein said polypeptide is derived from the species *Staphylococcus aureus*.
8. An antigenic polypeptide according to Claim 6 wherein said polypeptide is derived from the species *Staphylococcus epidermidis*.
9. An antigenic polypeptide according to any of Claims 1-8 wherein said polypeptide is an opsonin.
10. A vaccine composition comprising at least one antigenic polypeptide according to any of Claims 1-9.
11. A vaccine composition according to Claim 10 wherein said composition further comprises a carrier and/or an adjuvant.
12. A method to immunize an animal against a disease or condition caused by a pathogenic microbe comprising administering to said animal at least one antigenic polypeptide according to any of Claims 1-9 or a vaccine composition according to Claim 10 or 11.
13. A method according to Claim 12 wherein said animal is human.
14. A method according to Claim 12 or 13 wherein said disease or condition is selected from the group consisting of: bacterimia; septic shock; organ infection; skin infection; bacterial nasal colonisation; bacterial eye infections; septicaemia; tuberculosis; bacteria-associated food poisoning; blood infections; peritonitis; endocarditis; sepsis; meningitis; pneumonia; stomach ulcers; gonorrhoea; strep throat; streptococcal-associated toxic shock; necrotizing fasciitis; impetigo;

histoplasmosis; Lyme disease; gastro-enteritis; dysentery; shigellosis; *Staphylococcus aureus*-associated septicaemia, food-poisoning or skin disorders; *Staphylococcus epidermidis*-associated septicaemia, peritonitis or endocarditis.

5 15. A method according to Claim 14 wherein said disease or condition is the result of a *Staphylococcus spp* infection.

16. A method according to Claim 15 wherein said disease or condition is *Staphylococcus aureus*-associated septicaemia, food-poisoning or skin disorders.

10

17. A method according to Claim 15 wherein said disease or condition is *Staphylococcus epidermidis*-associated septicaemia, peritonitis or endocarditis.

18. An antibody, or binding part thereof, obtainable by the method according to
15 any of Claims 12-17.

19. An antibody according to Claim 18 wherein said antibody is a monoclonal antibody.

20 20. An antibody according to Claim 18 or 19 wherein said antibody is a chimeric antibody.

21. An antibody according to Claim 18 or 19 wherein said antibody is a humanized antibody.

25

22. An antibody according to any of Claims 18-21 wherein said antibody is an opsonic antibody.

23. An antibody according to any of Claims 18-22 wherein said antibody is a
30 therapeutic antibody or a diagnostic antibody.

24. A method for preparing a hybridoma cell-line producing monoclonal antibodies according to Claim 19 comprising the steps of:

- 5 i) immunising an immunocompetent mammal with an immunogen comprising at least one polypeptide having the amino acid sequence as represented in Tables 8 or 10, or polypeptide fragments thereof;
- ii) fusing lymphocytes of the immunised immunocompetent mammal with myeloma cells to form hybridoma cells;
- iii) screening monoclonal antibodies produced by the hybridoma cells of step (ii) for binding activity to the amino acid sequences of (i);
- 10 iv) culturing the hybridoma cells to proliferate and/or to secrete said monoclonal antibody; and optionally
- v) recovering the monoclonal antibody from the culture supernatant.

25. A method according to Claim 24 wherein said hybridoma cell-line produces
15 opsonic antibodies.

26. A hybridoma cell-line produced by the method of Claim 24 or 25.

27. A method to identify opsonic antigens expressed by a pathogenic microbe
20 comprising:

- i) providing a host cell transformed with a DNA library encoding genes, or partial gene sequences, of a pathogenic microbe;
- ii) providing conditions conducive to the expression of said transformed genes or partial sequences;
- 25 iii) contacting the antigens expressed by said gene sequences with autologous antisera derived from an animal infected with, or has been infected with, said pathogenic microbe;
- iv) purifying the DNA encoding antigenic polypeptides binding to said autologous antisera; and
- 30 v) testing the opsonic activity of a polypeptide encoded by said DNA molecule.